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(57) Abstract <p>Compounds and methods for treating lung cancer are provided. The inventive compounds include polypeptides containing at least a portion of a lung tumor protein. Vaccines and pharmaceutical compositions for immunotherapy of lung cancer comprising such polypeptides, or polynucleotides encoding such polypeptides, are also provided, together with polynucleotides for preparing the inventive polypeptides.</p>			

COMPOUNDS FOR THERAPY AND DIAGNOSIS OF LUNG CANCER AND METHODS FOR THEIR USE

5 TECHNICAL FIELD

The present invention relates generally to compositions and methods for the treatment of lung cancer. The invention is more specifically related to nucleotide sequences that are preferentially expressed in lung tumor tissue, together with polypeptides encoded by such nucleotide sequences. The inventive nucleotide sequences and polypeptides may be used
10 in vaccines and pharmaceutical compositions for the treatment of lung cancer.

BACKGROUND OF THE INVENTION

Lung cancer is the primary cause of cancer death among both men and women in the U.S., with an estimated 172,000 new cases being reported in 1994. The five-year
15 survival rate among all lung cancer patients, regardless of the stage of disease at diagnosis, is only 13%. This contrasts with a five-year survival rate of 46% among cases detected while the disease is still localized. However, only 16% of lung cancers are discovered before the disease has spread.

Early detection is difficult since clinical symptoms are often not seen until the
20 disease has reached an advanced stage. Currently, diagnosis is aided by the use of chest x-rays, analysis of the type of cells contained in sputum and fiberoptic examination of the bronchial passages. Treatment regimens are determined by the type and stage of the cancer, and include surgery, radiation therapy and/or chemotherapy. In spite of considerable research into therapies for the disease, lung cancer remains difficult to treat.

25 Accordingly, there remains a need in the art for improved vaccines, treatment methods and diagnostic techniques for lung cancer.

SUMMARY OF THE INVENTION

Briefly stated, the present invention provides compounds and methods for the
30 therapy of lung cancer. In a first aspect, isolated polynucleotides encoding lung tumor polypeptides are provided, such polynucleotides comprising a nucleotide sequence selected

herein; and (b) detecting in the sample a protein or polypeptide that binds to the binding agent. In preferred embodiments, the binding agent is an antibody, most preferably a monoclonal antibody.

In related aspects, methods are provided for monitoring the progression of lung cancer in a patient, comprising: (a) contacting a biological sample obtained from a patient with a binding agent that is capable of binding to one of the polypeptides disclosed herein; (b) determining in the sample an amount of a protein or polypeptide that binds to the binding agent; (c) repeating steps (a) and (b); and comparing the amounts of polypeptide detected in steps (b) and (c).

Within related aspects, the present invention provides antibodies, preferably monoclonal antibodies, that bind to the inventive polypeptides, as well as diagnostic kits comprising such antibodies, and methods of using such antibodies to inhibit the development of lung cancer.

The present invention further provides methods for detecting lung cancer comprising: (a) obtaining a biological sample from a patient; (b) contacting the sample with a first and a second oligonucleotide primer in a polymerase chain reaction, at least one of the oligonucleotide primers being specific for a polynucleotide that encodes one of the polypeptides disclosed herein; and (c) detecting in the sample a DNA sequence that amplifies in the presence of the first and second oligonucleotide primers. In a preferred embodiment, at least one of the oligonucleotide primers comprises at least about 10 contiguous nucleotides of a polynucleotide comprising a sequence selected from the group consisting of SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181.

In a further aspect, the present invention provides a method for detecting lung cancer in a patient comprising: (a) obtaining a biological sample from the patient; (b) contacting the sample with an oligonucleotide probe specific for a polynucleotide that encodes one of the polypeptides disclosed herein; and (c) detecting in the sample a DNA sequence that hybridizes to the oligonucleotide probe. Preferably, the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide comprising a sequence selected from the group consisting of SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181. In related aspects, diagnostic kits comprising the above oligonucleotide probes or primers are provided.

- SEQ ID NO: 14 is the determined cDNA sequence for L355C1.cons
SEQ ID NO: 15 is the determined cDNA sequence for L366C1.cons
SEQ ID NO: 16 is the determined cDNA sequence for L163C1a
SEQ ID NO: 17 is the determined cDNA sequence for LT86-1
5 SEQ ID NO: 18 is the determined cDNA sequence for LT86-2
SEQ ID NO: 19 is the determined cDNA sequence for LT86-3
SEQ ID NO: 20 is the determined cDNA sequence for LT86-4
SEQ ID NO: 21 is the determined cDNA sequence for LT86-5
SEQ ID NO: 22 is the determined cDNA sequence for LT86-6
10 SEQ ID NO: 23 is the determined cDNA sequence for LT86-7
SEQ ID NO: 24 is the determined cDNA sequence for LT86-8
SEQ ID NO: 25 is the determined cDNA sequence for LT86-9
SEQ ID NO: 26 is the determined cDNA sequence for LT86-10
SEQ ID NO: 27 is the determined cDNA sequence for LT86-11
15 SEQ ID NO: 28 is the determined cDNA sequence for LT86-12
SEQ ID NO: 29 is the determined cDNA sequence for LT86-13
SEQ ID NO: 30 is the determined cDNA sequence for LT86-14
SEQ ID NO: 31 is the determined cDNA sequence for LT86-15
SEQ ID NO: 32 is the predicted amino acid sequence for LT86-1
20 SEQ ID NO: 33 is the predicted amino acid sequence for LT86-2
SEQ ID NO: 34 is the predicted amino acid sequence for LT86-3
SEQ ID NO: 35 is the predicted amino acid sequence for LT86-4
SEQ ID NO: 36 is the predicted amino acid sequence for LT86-5
SEQ ID NO: 37 is the predicted amino acid sequence for LT86-6
25 SEQ ID NO: 38 is the predicted amino acid sequence for LT86-7
SEQ ID NO: 39 is the predicted amino acid sequence for LT86-8
SEQ ID NO: 40 is the predicted amino acid sequence for LT86-9
SEQ ID NO: 41 is the predicted amino acid sequence for LT86-10
SEQ ID NO: 42 is the predicted amino acid sequence for LT86-11
30 SEQ ID NO: 43 is the predicted amino acid sequence for LT86-12

- SEQ ID NO: 74 is the predicted amino acid sequence for LT86-21
SEQ ID NO: 75 is the predicted amino acid sequence for LT86-22
SEQ ID NO: 76 is the predicted amino acid sequence for LT86-26
SEQ ID NO: 77 is the predicted amino acid sequence for LT86-27
5 SEQ ID NO: 78 is the determined extended cDNA sequence for L86S-12
SEQ ID NO: 79 is the determined extended cDNA sequence for L86S-36
SEQ ID NO: 80 is the determined extended cDNA sequence for L86S-46
SEQ ID NO: 81 is the predicted extended amino acid sequence for L86S-12
SEQ ID NO: 82 is the predicted extended amino acid sequence for L86S-36
10 SEQ ID NO: 83 is the predicted extended amino acid sequence for L86S-46
SEQ ID NO: 84 is the determined 5'cDNA sequence for L86S-6
SEQ ID NO: 85 is the determined 5'cDNA sequence for L86S-11
SEQ ID NO: 86 is the determined 5'cDNA sequence for L86S-14
SEQ ID NO: 87 is the determined 5'cDNA sequence for L86S-29
15 SEQ ID NO: 88 is the determined 5'cDNA sequence for L86S-34
SEQ ID NO: 89 is the determined 5'cDNA sequence for L86S-39
SEQ ID NO: 90 is the determined 5'cDNA sequence for L86S-47
SEQ ID NO: 91 is the determined 5'cDNA sequence for L86S-49
SEQ ID NO: 92 is the determined 5'cDNA sequence for L86S-51
20 SEQ ID NO: 93 is the predicted amino acid sequence for L86S-6
SEQ ID NO: 94 is the predicted amino acid sequence for L86S-11
SEQ ID NO: 95 is the predicted amino acid sequence for L86S-14
SEQ ID NO: 96 is the predicted amino acid sequence for L86S-29
SEQ ID NO: 97 is the predicted amino acid sequence for L86S-34
25 SEQ ID NO: 98 is the predicted amino acid sequence for L86S-39
SEQ ID NO: 99 is the predicted amino acid sequence for L86S-47
SEQ ID NO: 100 is the predicted amino acid sequence for L86S-49
SEQ ID NO: 101 is the predicted amino acid sequence for L86S-51
SEQ ID NO: 102 is the determined DNA sequence for SLT-T1
30 SEQ ID NO: 103 is the determined 5' cDNA sequence for SLT-T2

- SEQ ID NO: 134 is the determined cDNA sequence for PSLT-69
SEQ ID NO: 135 is the determined cDNA sequence for PSLT-71
SEQ ID NO: 136 is the determined cDNA sequence for PSLT-73
SEQ ID NO: 137 is the determined cDNA sequence for PSLT-79
5 SEQ ID NO: 138 is the determined cDNA sequence for PSLT-03
SEQ ID NO: 139 is the determined cDNA sequence for PSLT-09
SEQ ID NO: 140 is the determined cDNA sequence for PSLT-011
SEQ ID NO: 141 is the determined cDNA sequence for PSLT-041
SEQ ID NO: 142 is the determined cDNA sequence for PSLT-62
10 SEQ ID NO: 143 is the determined cDNA sequence for PSLT-6
SEQ ID NO: 144 is the determined cDNA sequence for PSLT-37
SEQ ID NO: 145 is the determined cDNA sequence for PSLT-74
SEQ ID NO: 146 is the determined cDNA sequence for PSLT-010
SEQ ID NO: 147 is the determined cDNA sequence for PSLT-012
15 SEQ ID NO: 148 is the determined cDNA sequence for PSLT-037
SEQ ID NO: 149 is the determined 5' cDNA sequence for SAL-3
SEQ ID NO: 150 is the determined 5' cDNA sequence for SAL-24
SEQ ID NO: 151 is the determined 5' cDNA sequence for SAL-25
SEQ ID NO: 152 is the determined 5' cDNA sequence for SAL-33
20 SEQ ID NO: 153 is the determined 5' cDNA sequence for SAL-50
SEQ ID NO: 154 is the determined 5' cDNA sequence for SAL-57
SEQ ID NO: 155 is the determined 5' cDNA sequence for SAL-66
SEQ ID NO: 156 is the determined 5' cDNA sequence for SAL-82
SEQ ID NO: 157 is the determined 5' cDNA sequence for SAL-99
25 SEQ ID NO: 158 is the determined 5' cDNA sequence for SAL-104
SEQ ID NO: 159 is the determined 5' cDNA sequence for SAL-109
SEQ ID NO: 160 is the determined 5' cDNA sequence for SAL-5
SEQ ID NO: 161 is the determined 5' cDNA sequence for SAL-8
SEQ ID NO: 162 is the determined 5' cDNA sequence for SAL-12
30 SEQ ID NO: 163 is the determined 5' cDNA sequence for SAL-14

- SEQ ID NO: 194 is the predicted amino acid sequence for SAL-5
SEQ ID NO: 195 is the predicted amino acid sequence for SAL-8
SEQ ID NO: 196 is the predicted amino acid sequence for SAL-12
SEQ ID NO: 197 is the predicted amino acid sequence for SAL-14
5 SEQ ID NO: 198 is the predicted amino acid sequence for SAL-16
SEQ ID NO: 199 is the predicted amino acid sequence for SAL-23
SEQ ID NO: 200 is the predicted amino acid sequence for SAL-26
SEQ ID NO: 201 is the predicted amino acid sequence for SAL-29
SEQ ID NO: 202 is the predicted amino acid sequence for SAL-32
10 SEQ ID NO: 203 is the predicted amino acid sequence for SAL-39
SEQ ID NO: 204 is the predicted amino acid sequence for SAL-42
SEQ ID NO: 205 is the predicted amino acid sequence for SAL-43
SEQ ID NO: 206 is the predicted amino acid sequence for SAL-44
SEQ ID NO: 207 is the predicted amino acid sequence for SAL-48
15 SEQ ID NO: 208 is the predicted amino acid sequence for SAL-68
SEQ ID NO: 209 is the predicted amino acid sequence for SAL-72
SEQ ID NO: 210 is the predicted amino acid sequence for SAL-77
SEQ ID NO: 211 is the predicted amino acid sequence for SAL-86
SEQ ID NO: 212 is the predicted amino acid sequence for SAL-88
20 SEQ ID NO: 213 is the predicted amino acid sequence for SAL-93
SEQ ID NO: 214 is the predicted amino acid sequence for SAL-100
SEQ ID NO: 215 is the predicted amino acid sequence for SAL-105
SEQ ID NO: 216 is a second predicted amino acid sequence for SAL-50

25 DETAILED DESCRIPTION OF THE INVENTION

As noted above, the present invention is generally directed to compositions and methods for the therapy of lung cancer. The compositions described herein include polypeptides, fusion proteins and polynucleotides. Also included within the present invention are molecules (such as an antibody or fragment thereof) that bind to the inventive
30 polypeptides. Such molecules are referred to herein as "binding agents."

of the proteins described herein may be identified in antibody binding assays. Such assays may generally be performed using any of a variety of means known to those of ordinary skill in the art, as described, for example, in Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, 1988. For example, a polypeptide
5 may be immobilized on a solid support (as described below) and contacted with patient sera to allow binding of antibodies within the sera to the immobilized polypeptide. Unbound sera may then be removed and bound antibodies detected using, for example, ¹²⁵I-labeled Protein A. Alternatively, a polypeptide may be used to generate monoclonal and polyclonal antibodies for use in detection of the polypeptide in blood or other fluids of lung cancer
10 patients. Methods for preparing and identifying immunogenic portions of antigens of known sequence are well known in the art and include those summarized in Paul, *Fundamental Immunology*, 3rd ed., Raven Press, 1993, pp. 243-247.

The term "polynucleotide(s)," as used herein, means a single or double-stranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and
15 corresponding RNA molecules, including HnRNA and mRNA molecules, both sense and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA, as well as wholly or partially synthesized polynucleotides. An HnRNA molecule contains introns and corresponds to a DNA molecule in a generally one-to-one manner. An mRNA molecule corresponds to an HnRNA and DNA molecule from which the introns have been excised. A
20 polynucleotide may consist of an entire gene, or any portion thereof. Operable anti-sense polynucleotides may comprise a fragment of the corresponding polynucleotide, and the definition of "polynucleotide" therefore includes all such operable anti-sense fragments.

The compositions and methods of the present invention also encompass variants of the above polypeptides and polynucleotides.

25 A polypeptide "variant," as used herein, is a polypeptide that differs from the recited polypeptide only in conservative substitutions and/or modifications, such that the antigenic properties of the polypeptide are retained. In a preferred embodiment, variant polypeptides differ from an identified sequence by substitution, deletion or addition of five amino acids or fewer. Such variants may generally be identified by modifying one of the
30 above polypeptide sequences, and evaluating the antigenic properties of the modified polypeptide using, for example, the representative procedures described herein. Polypeptide

SSC, 0.5% SDS, 1.0 mM EDTA (pH 8.0); hybridizing at 50°C–65°C, 5X SSC, overnight or, in the event of cross-species homology, at 45°C with 0.5X SSC; followed by washing twice at 65°C for 20 minutes with each of 2X, 0.5X and 0.2X SSC containing 0.1% SDS. Such hybridizing DNA sequences are also within the scope of this invention, as are nucleotide
5 sequences that, due to code degeneracy, encode an immunogenic polypeptide that is encoded by a hybridizing DNA sequence.

Two nucleotide or polypeptide sequences are said to be “identical” if the sequence of nucleotides or amino acid residues in the two sequences is the same when aligned for maximum correspondence as described below. Comparisons between two sequences are
10 typically performed by comparing the sequences over a comparison window to identify and compare local regions of sequence similarity. A “comparison window” as used herein, refers to a segment of at least about 20 contiguous positions, usually 30 to about 75, 40 to about 50, in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned.

15 Optimal alignment of sequences for comparison may be conducted using the Megalign program in the Lasergene suite of bioinformatics software (DNASTAR, Inc., Madison, WI), using default parameters. This program embodies several alignment schemes described in the following references: Dayhoff, M.O. (1978) A model of evolutionary change in proteins – Matrices for detecting distant relationships. In Dayhoff, M.O. (ed.) Atlas of
20 Protein Sequence and Structure, National Biomedical Research Foundation, Washington DC Vol. 5, Suppl. 3, pp. 345–358; Hein J. (1990) Unified Approach to Alignment and Phylogenesis pp. 626–645 *Methods in Enzymology* vol. 183, Academic Press, Inc., San Diego, CA; Higgins, D.G. and Sharp, P.M. (1989) Fast and sensitive multiple sequence alignments on a microcomputer *CABIOS* 5:151–153; Myers, E.W. and Muller W. (1988) Optimal alignments
25 in linear space *CABIOS* 4:11–17; Robinson, E.D. (1971) *Comb. Theor* 11:105; Santou, N. Nes, M. (1987) The neighbor joining method. A new method for reconstructing phylogenetic trees *Mol. Biol. Evol.* 4:406–425; Sneath, P.H.A. and Sokal, R.R. (1973) *Numerical Taxonomy – the Principles and Practice of Numerical Taxonomy*, Freeman Press, San Francisco, CA; Wilbur, W.J. and Lipman, D.J. (1983) Rapid similarity searches of nucleic
30 acid and protein data banks *Proc. Natl. Acad. Sci. USA* 80:726–730.

libraries prepared from SCID mice with mouse anti-tumor sera, as described below in Example 4. Examples of cDNA sequences that may be isolated using this technique are provided in SEQ ID NO: 149-181.

5 A gene encoding a polypeptide described herein (or a portion thereof) may, alternatively, be amplified from human genomic DNA, or from lung tumor cDNA, via polymerase chain reaction. For this approach, sequence-specific primers may be designed based on the nucleotide sequences provided herein and may be purchased or synthesized. An amplified portion of a specific nucleotide sequence may then be used to isolate the full length gene from a human genomic DNA library or from a lung tumor cDNA library, using well
10 known techniques, such as those described in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratories, Cold Spring Harbor, NY (1989).

Once a DNA sequence encoding a polypeptide is obtained, the polypeptide may be produced recombinantly by inserting the DNA sequence into an expression vector and expressing the polypeptide in an appropriate host. Any of a variety of expression vectors
15 known to those of ordinary skill in the art may be employed to express recombinant polypeptides of this invention. Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polynucleotide that encodes the recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are *E. coli*, yeast or a mammalian
20 cell line, such as COS or CHO cells. The DNA sequences expressed in this manner may encode naturally occurring polypeptides, portions of naturally occurring polypeptides, or other variants thereof. Supernatants from suitable host/vector systems which secrete the recombinant polypeptide may be first concentrated using a commercially available filter. The concentrate may then be applied to a suitable purification matrix, such as an affinity matrix or
25 ion exchange resin. Finally, one or more reverse phase HPLC steps can be employed to further purify the recombinant polypeptide.

Such techniques may also be used to prepare polypeptides comprising portions or variants of the native polypeptides. Portions and other variants having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using
30 techniques well known to those of ordinary skill in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as

extended conformation; (2) their inability to adopt a secondary structure that could interact with functional epitopes on the first and second polypeptides; and (3) the lack of hydrophobic or charged residues that might react with the polypeptide functional epitopes. Preferred peptide linker sequences contain Gly, Asn and Ser residues. Other near neutral amino acids, such as Thr and Ala may also be used in the linker sequence. Amino acid sequences which may be usefully employed as linkers include those disclosed in Maratea et al., *Gene* 40:39-46, 1985; Murphy et al., *Proc. Natl. Acad. Sci. USA* 83:8258-8262, 1986; U.S. Patent No. 4,935,233 and U.S. Patent No. 4,751,180. The linker sequence may be from 1 to about 50 amino acids in length. Peptide sequences are not required when the first and second polypeptides have non-essential N-terminal amino acid regions that can be used to separate the functional domains and prevent steric interference.

The ligated DNA sequences are operably linked to suitable transcriptional or translational regulatory elements. The regulatory elements responsible for expression of DNA are located only 5' to the DNA sequence encoding the first polypeptides. Similarly, stop codons require to end translation and transcription termination signals are only present 3' to the DNA sequence encoding the second polypeptide.

Fusion proteins are also provided that comprise a polypeptide of the present invention together with an unrelated immunogenic protein. Preferably the immunogenic protein is capable of eliciting a recall response. Examples of such proteins include tetanus, tuberculosis and hepatitis proteins (see, for example, Stoute et al. *New Engl. J. Med.*, 336:86-91 (1997)).

Polypeptides that comprise an immunogenic portion of a lung tumor protein may generally be used for therapy of lung cancer, wherein the polypeptide stimulates the patient's own immune response to lung tumor cells. The present invention thus provides methods for using one or more of the compounds described herein (which may be polypeptides, polynucleotides or fusion proteins) for immunotherapy of lung cancer in a patient. As used herein, a "patient" refers to any warm-blooded animal, preferably a human. A patient may be afflicted with disease, or may be free of detectable disease. Accordingly, the compounds disclosed herein may be used to treat lung cancer or to inhibit the development of lung cancer. In a preferred embodiment, the compounds are administered

ordinary skill in the art. The DNA may also be "naked," as described, for example, in published PCT application WO 90/11092, and Ulmer et al., *Science* 259:1745-1749, 1993, reviewed by Cohen, *Science* 259:1691-1692, 1993. The uptake of naked DNA may be increased by coating the DNA onto biodegradable beads, which are efficiently transported into the cells.

Routes and frequency of administration, as well as dosage, will vary from individual to individual and may parallel those currently being used in immunotherapy of other diseases. In general, the pharmaceutical compositions and vaccines may be administered by injection (e.g., intracutaneous, intramuscular, intravenous or subcutaneous), intranasally (e.g., by aspiration) or orally. Between 1 and 10 doses may be administered over a 3-24 week period. Preferably, 4 doses are administered, at an interval of 3 months, and booster administrations may be given periodically thereafter. Alternate protocols may be appropriate for individual patients. A suitable dose is an amount of polypeptide or DNA that is effective to raise an immune response (cellular and/or humoral) against lung tumor cells in a treated patient. A suitable immune response is at least 10-50% above the basal (i.e., untreated) level. In general, the amount of polypeptide present in a dose (or produced *in situ* by the DNA in a dose) ranges from about 1 pg to about 100 mg per kg of host, typically from about 10 pg to about 1 mg, and preferably from about 100 pg to about 1 μ g. Suitable dose sizes will vary with the size of the patient, but will typically range from about 0.01 mL to about 5 mL.

While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this invention, the type of carrier will vary depending on the mode of administration. For parenteral administration, such as subcutaneous injection, the carrier preferably comprises water, saline, alcohol, a lipid, a wax and/or a buffer. For oral administration, any of the above carriers or a solid carrier, such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, and/or magnesium carbonate, may be employed. Biodegradable microspheres (e.g., polylactic glycolide) may also be employed as carriers for the pharmaceutical compositions of this invention. Suitable biodegradable microspheres are disclosed, for example, in U.S. Patent Nos. 4,897,268 and 5,075,109.

(Natural Killer cells, lymphokine-activated killer cells), B cells, or antigen presenting cells (such as dendritic cells and macrophages) expressing the disclosed antigens. The polypeptides disclosed herein may also be used to generate antibodies or anti-idiotypic antibodies (as in U.S. Patent No. 4,918,164), for passive immunotherapy.

5 The predominant method of procuring adequate numbers of T-cells for adoptive immunotherapy is to grow immune T-cells *in vitro*. Culture conditions for expanding single antigen-specific T-cells to several billion in number with retention of antigen recognition *in vivo* are well known in the art. These *in vitro* culture conditions typically utilize intermittent stimulation with antigen, often in the presence of cytokines, such
10 as IL-2, and non-dividing feeder cells. As noted above, the immunoreactive polypeptides described herein may be used to rapidly expand antigen-specific T cell cultures in order to generate sufficient number of cells for immunotherapy. In particular, antigen-presenting cells, such as dendritic, macrophage or B-cells, may be pulsed with immunoreactive polypeptides or transfected with a polynucleotide sequence(s), using standard techniques well
15 known in the art. For cultured T-cells to be effective in therapy, the cultured T-cells must be able to grow and distribute widely and to survive long term *in vivo*. Studies have demonstrated that cultured T-cells can be induced to grow *in vivo* and to survive long term in substantial numbers by repeated stimulation with antigen supplemented with IL-2 (see, for example, Cheever et al. *Ibid*).

20 The polypeptides disclosed herein may also be employed to generate and/or isolate tumor-reactive T-cells, which can then be administered to the patient. In one technique, antigen-specific T-cell lines may be generated by *in vivo* immunization with short peptides corresponding to immunogenic portions of the disclosed polypeptides. The resulting antigen specific CD8+ CTL clones may be isolated from the patient, expanded using standard
25 tissue culture techniques, and returned to the patient.

 Alternatively, peptides corresponding to immunogenic portions of the polypeptides may be employed to generate tumor reactive T cell subsets by selective *in vitro* stimulation and expansion of autologous T cells to provide antigen-specific T cells which may be subsequently transferred to the patient as described, for example, by Chang et al.
30 (*Crit. Rev. Oncol. Hematol.*, 22(3), 213, 1996).

at least about 80%, and preferably at least about 90%) of the patients for which lung cancer would be indicated using the full length protein, and that indicate the absence of lung cancer in substantially all of those samples that would be negative when tested with full length protein. The representative assays described below, such as the two-antibody sandwich
5 assay, may generally be employed for evaluating the ability of a binding agent to detect metastatic human lung tumors.

The ability of a polypeptide prepared as described herein to generate antibodies capable of detecting primary or metastatic human lung tumors may generally be evaluated by raising one or more antibodies against the polypeptide (using, for example, a
10 representative method described herein) and determining the ability of such antibodies to detect such tumors in patients. This determination may be made by assaying biological samples from patients with and without primary or metastatic lung cancer for the presence of a polypeptide that binds to the generated antibodies. Such test assays may be performed, for example, using a representative procedure described below. Polypeptides that generate
15 antibodies capable of detecting at least 20% of primary or metastatic lung tumors by such procedures are considered to be useful in assays for detecting primary or metastatic human lung tumors. Polypeptide specific antibodies may be used alone or in combination to improve sensitivity.

Polypeptides capable of detecting primary or metastatic human lung tumors
20 may be used as markers for diagnosing lung cancer or for monitoring disease progression in patients. In one embodiment, lung cancer in a patient may be diagnosed by evaluating a biological sample obtained from the patient for the level of one or more of the above polypeptides, relative to a predetermined cut-off value. As used herein, suitable "biological samples" include blood, sera, urine and/or lung secretions.

25 The level of one or more of the above polypeptides may be evaluated using any binding agent specific for the polypeptide(s). A "binding agent," in the context of this invention, is any agent (such as a compound or a cell) that binds to a polypeptide as described above. As used herein, "binding" refers to a noncovalent association between two separate molecules (each of which may be free (*i.e.*, in solution) or present on the surface of a cell or a
30 solid support), such that a "complex" is formed. Such a complex may be free or immobilized (either covalently or noncovalently) on a support material. The ability to bind may generally

be immobilized on the solid support using a variety of techniques known to those of skill in the art, which are amply described in the patent and scientific literature. In the context of the present invention, the term "immobilization" refers to both noncovalent association, such as adsorption, and covalent attachment (which may be a direct linkage between the antigen and functional groups on the support or may be a linkage by way of a cross-linking agent).
5 Immobilization by adsorption to a well in a microtiter plate or to a membrane is preferred. In such cases, adsorption may be achieved by contacting the binding agent, in a suitable buffer, with the solid support for a suitable amount of time. The contact time varies with temperature, but is typically between about 1 hour and about 1 day. In general, contacting a well of a plastic microtiter plate (such as polystyrene or polyvinylchloride) with an amount of
10 binding agent ranging from about 10 ng to about 10 μ g, and preferably about 100 ng to about 1 μ g, is sufficient to immobilize an adequate amount of binding agent.

Covalent attachment of binding agent to a solid support may generally be achieved by first reacting the support with a bifunctional reagent that will react with both the support and a functional group, such as a hydroxyl or amino group, on the binding agent. For
15 example, the binding agent may be covalently attached to supports having an appropriate polymer coating using benzoquinone or by condensation of an aldehyde group on the support with an amine and an active hydrogen on the binding partner (*see, e.g.,* Pierce Immunotechnology Catalog and Handbook, 1991, at A12-A13).

20 In certain embodiments, the assay is a two-antibody sandwich assay. This assay may be performed by first contacting an antibody that has been immobilized on a solid support, commonly the well of a microtiter plate, with the sample, such that polypeptides within the sample are allowed to bind to the immobilized antibody. Unbound sample is then removed from the immobilized polypeptide-antibody complexes and a second antibody
25 (containing a reporter group) capable of binding to a different site on the polypeptide is added. The amount of second antibody that remains bound to the solid support is then determined using a method appropriate for the specific reporter group.

More specifically, once the antibody is immobilized on the support as described above, the remaining protein binding sites on the support are typically blocked.
30 Any suitable blocking agent known to those of ordinary skill in the art, such as bovine serum albumin or Tween 20™ (Sigma Chemical Co., St. Louis, MO). The immobilized antibody is

that corresponds to a predetermined cut-off value. In one preferred embodiment, the cut-off value is the average mean signal obtained when the immobilized antibody is incubated with samples from patients without lung cancer. In general, a sample generating a signal that is three standard deviations above the predetermined cut-off value is considered positive for lung cancer. In an alternate preferred embodiment, the cut-off value is determined using a Receiver Operator Curve, according to the method of Sackett et al., *Clinical Epidemiology: A Basic Science for Clinical Medicine*, Little Brown and Co., 1985, p. 106-7. Briefly, in this embodiment, the cut-off value may be determined from a plot of pairs of true positive rates (i.e., sensitivity) and false positive rates (100%-specificity) that correspond to each possible cut-off value for the diagnostic test result. The cut-off value on the plot that is the closest to the upper left-hand corner (i.e., the value that encloses the largest area) is the most accurate cut-off value, and a sample generating a signal that is higher than the cut-off value determined by this method may be considered positive. Alternatively, the cut-off value may be shifted to the left along the plot, to minimize the false positive rate, or to the right, to minimize the false negative rate. In general, a sample generating a signal that is higher than the cut-off value determined by this method is considered positive for lung cancer.

In a related embodiment, the assay is performed in a flow-through or strip test format, wherein the antibody is immobilized on a membrane, such as nitrocellulose. In the flow-through test, polypeptides within the sample bind to the immobilized antibody as the sample passes through the membrane. A second, labeled antibody then binds to the antibody-polypeptide complex as a solution containing the second antibody flows through the membrane. The detection of bound second antibody may then be performed as described above. In the strip test format, one end of the membrane to which antibody is bound is immersed in a solution containing the sample. The sample migrates along the membrane through a region containing second antibody and to the area of immobilized antibody. Concentration of second antibody at the area of immobilized antibody indicates the presence of lung cancer. Typically, the concentration of second antibody at that site generates a pattern, such as a line, that can be read visually. The absence of such a pattern indicates a negative result. In general, the amount of antibody immobilized on the membrane is selected to generate a visually discernible pattern when the biological sample contains a level of polypeptide that would be sufficient to generate a positive signal in the two-antibody

of immortal cell lines capable of producing antibodies having the desired specificity (*i.e.*, reactivity with the polypeptide of interest). Such cell lines may be produced, for example, from spleen cells obtained from an animal immunized as described above. The spleen cells are then immortalized by, for example, fusion with a myeloma cell fusion partner, preferably one that is syngeneic with the immunized animal. A variety of fusion techniques may be employed. For example, the spleen cells and myeloma cells may be combined with a nonionic detergent for a few minutes and then plated at low density on a selective medium that supports the growth of hybrid cells, but not myeloma cells. A preferred selection technique uses HAT (hypoxanthine, aminopterin, thymidine) selection. After a sufficient time, usually about 1 to 2 weeks, colonies of hybrids are observed. Single colonies are selected and tested for binding activity against the polypeptide. Hybridomas having high reactivity and specificity are preferred.

Monoclonal antibodies may be isolated from the supernatants of growing hybridoma colonies. In addition, various techniques may be employed to enhance the yield, such as injection of the hybridoma cell line into the peritoneal cavity of a suitable vertebrate host, such as a mouse. Monoclonal antibodies may then be harvested from the ascites fluid or the blood. Contaminants may be removed from the antibodies by conventional techniques, such as chromatography, gel filtration, precipitation, and extraction. The polypeptides of this invention may be used in the purification process in, for example, an affinity chromatography step.

Monoclonal antibodies of the present invention may also be used as therapeutic reagents, to diminish or eliminate lung tumors. The antibodies may be used on their own (for instance, to inhibit metastases) or coupled to one or more therapeutic agents. Suitable agents in this regard include radionuclides, differentiation inducers, drugs, toxins, and derivatives thereof. Preferred radionuclides include ^{90}Y , ^{123}I , ^{125}I , ^{131}I , ^{186}Re , ^{188}Re , ^{211}At , and ^{212}Bi . Preferred drugs include methotrexate, and pyrimidine and purine analogs. Preferred differentiation inducers include phorbol esters and butyric acid. Preferred toxins include ricin, abrin, diphtheria toxin, cholera toxin, gelonin, *Pseudomonas* exotoxin, *Shigella* toxin, and pokeweed antiviral protein.

A therapeutic agent may be coupled (*e.g.*, covalently bonded) to a suitable monoclonal antibody either directly or indirectly (*e.g.*, via a linker group). A direct reaction

be prepared in a variety of ways. For example, more than one agent may be coupled directly to an antibody molecule, or linkers which provide multiple sites for attachment can be used. Alternatively, a carrier can be used.

A carrier may bear the agents in a variety of ways, including covalent bonding either directly or via a linker group. Suitable carriers include proteins such as albumins (*e.g.*, U.S. Patent No. 4,507,234, to Kato et al.), peptides and polysaccharides such as aminodextran (*e.g.*, U.S. Patent No. 4,699,784, to Shih et al.). A carrier may also bear an agent by noncovalent bonding or by encapsulation, such as within a liposome vesicle (*e.g.*, U.S. Patent Nos. 4,429,008 and 4,873,088). Carriers specific for radionuclide agents include radiohalogenated small molecules and chelating compounds. For example, U.S. Patent No. 4,735,792 discloses representative radiohalogenated small molecules and their synthesis. A radionuclide chelate may be formed from chelating compounds that include those containing nitrogen and sulfur atoms as the donor atoms for binding the metal, or metal oxide, radionuclide. For example, U.S. Patent No. 4,673,562, to Davison et al. discloses representative chelating compounds and their synthesis.

A variety of routes of administration for the antibodies and immunoconjugates may be used. Typically, administration will be intravenous, intramuscular, subcutaneous or in the bed of a resected tumor. It will be evident that the precise dose of the antibody/immunoconjugate will vary depending upon the antibody used, the antigen density on the tumor, and the rate of clearance of the antibody.

Diagnostic reagents of the present invention may also comprise DNA sequences encoding one or more of the above polypeptides, or one or more portions thereof. For example, at least two oligonucleotide primers may be employed in a polymerase chain reaction (PCR) based assay to amplify lung tumor-specific cDNA derived from a biological sample, wherein at least one of the oligonucleotide primers is specific for a polynucleotide encoding a lung tumor protein of the present invention. The presence of the amplified cDNA is then detected using techniques well known in the art, such as gel electrophoresis. Similarly, oligonucleotide probes specific for a polynucleotide encoding a lung tumor protein of the present invention may be used in a hybridization assay to detect the presence of an inventive polypeptide in a biological sample.

The following Examples are offered by way of illustration and not by way of limitation.

EXAMPLES

Example 1

PREPARATION OF LUNG TUMOR-SPECIFIC cDNA SEQUENCES USING DIFFERENTIAL DISPLAY RT-PCR

This example illustrates the preparation of cDNA molecules encoding lung tumor-specific polypeptides using a differential display screen.

Tissue samples were prepared from breast tumor and normal tissue of a patient with lung cancer that was confirmed by pathology after removal of samples from the patient. Normal RNA and tumor RNA was extracted from the samples and mRNA was isolated and converted into cDNA using a (dT)₁₂AG (SEQ ID NO: 47) anchored 3' primer. Differential display PCR was then executed using a randomly chosen primer (SEQ ID NO: 48). Amplification conditions were standard buffer containing 1.5 mM MgCl₂, 20 pmol of primer, 500 pmol dNTP and 1 unit of Taq DNA polymerase (Perkin-Elmer, Branchburg, NJ). Forty cycles of amplification were performed using 94 °C denaturation for 30 seconds, 42 °C annealing for 1 minute and 72 °C extension for 30 seconds. Bands that were repeatedly observed to be specific to the RNA fingerprint pattern of the tumor were cut out of a silver stained gel, subcloned into the pGEM-T vector (Promega, Madison, WI) and sequenced. The isolated 3' sequences are provided in SEQ ID NO: 1-16.

Comparison of these sequences to those in the public databases using the BLASTN program, revealed no significant homologies to the sequences provided in SEQ ID NO: 1-11. To the best of the inventors' knowledge, none of the isolated DNA sequences have previously been shown to be expressed at a greater level in human lung tumor tissue than in normal lung tissue.

aminopeptidase. Clone LT86-9 appears to contain two inserts, with the 5' sequence showing homology to the previously identified antisense sequence of interferon alpha-induced P27, and the 3' sequence being similar to LT86-6. Clone LT86-14 (SEQ ID NO: 30) was found to show some homology to the trithorax gene and has an "RGD" cell attachment sequence and a
5 beta-Lactamase A site which functions in hydrolysis of penicillin. Clones LT86-1, LT86-2, LT86-4, LT86-5 and LT86-10 (SEQ ID NOS: 17, 18, 20, 21 and 26, respectively) were found to show homology to previously identified genes. A subsequently determined extended cDNA sequence for LT86-4 is provided in SEQ ID NO: 66, with the corresponding predicted amino acid sequence being provided in SEQ ID NO: 67.

10 Subsequent studies led to the isolation of five additional clones, referred to as LT86-20, LT86-21, LT86-22, LT86-26 and LT86-27. The determined 5' cDNA sequences for LT86-20, LT86-22, LT86-26 and LT86-27 are provided in SEQ ID NO: 68 and 70-72, respectively, with the determined 3' cDNA sequences for LT86-21 being provided in SEQ ID
15 NO: 69. The corresponding predicted amino acid sequences for LT86-20, LT86-21, LT86-22, LT86-26 and LT86-27 are provided in SEQ ID NO: 73-77, respectively. LT86-22 and LT86-27 were found to be highly similar to each other. Comparison of these sequences to those in the gene bank as described above, revealed no significant homologies to LT86-22 and LT86-27. LT86-20, LT86-21 and LT86-26 were found to show homology to previously
20 identified genes.

predicted amino acid sequences are provided in SEQ ID NO: 93-101, respectively. L86S-30, L86S-39 and L86S-47 were found to be similar to each other. Comparison of these sequences with those in the gene bank as described above, revealed no significant homologies to L86S-14. L86S-29 was found to show some homology to a previously identified EST. 5 L86S-6, L86S-11, L86S-34, L86S-39, L86S-47, L86S-49 and L86S-51 were found to show some homology to previously identified genes.

In further studies, a directional cDNA library was constructed using a Stratagene kit with a Lambda Zap Express vector. Total RNA for the library was isolated from two primary squamous lung tumors and poly A+ RNA was isolated using an oligo dT 10 column. Antiserum was developed in normal mice using a pool of sera from three SCID mice implanted with human squamous lung carcinomas. Approximately 700,000 PFUs were screened from the unamplified library with *E. coli* absorbed mouse anti-SCID tumor serum. Positive plaques were identified as described above. Phage was purified and phagemid excised for 180 clones with inserts in a pBK-CMV vector for expression in prokaryotic or 15 eukaryotic cells.

The determined cDNA sequences for 23 of the isolated clones are provided in SEQ ID NO: 126-148. Comparison of these sequences with those in the public database as described above revealed no significant homologies to the sequences of SEQ ID NO: 139 and 143-148. The sequences of SEQ ID NO: 126-138 and 140-142 were found to show 20 homology previously identified human polynucleotide sequences.

tags (ESTs). The sequences of SEQ ID NO: 150, 155 and 159-181 were found to show homology to sequences previously identified in humans.

Example 6

ISOLATION OF DNA SEQUENCES ENCODING LUNG TUMOR ANTIGENS

DNA sequences encoding antigens potentially involved in squamous cell lung
5 tumor formation were isolated as follows.

A lung tumor directional cDNA expression library was constructed employing
the Lambda ZAP Express expression system (Stratagene, La Jolla, CA). Total RNA for the
library was taken from a pool of two human squamous epithelial lung carcinomas and poly
A+ RNA was isolated using oligo-dT cellulose (Gibco BRL, Gaithersburg, MD). Phagemid
10 were rescued at random and the cDNA sequences of isolated clones were determined.

The determined cDNA sequence for the clone SLT-T1 is provided in SEQ ID
NO: 102, with the determined 5' cDNA sequences for the clones SLT-T2, SLT-T3, SLT-T5,
SLT-T7, SLT-T9, SLT-T10, SLT-T11 and SLT-T12 being provided in SEQ ID NO: 103-
110, respectively. The corresponding predicted amino acid sequence for SLT-T1, SLT-T2,
15 SLT-T3, SLT-T10 and SLT-T12 are provided in SEQ ID NO: 111-115, respectively.
Comparison of the sequences for SLT-T2, SLT-T3, SLT-T5, SLT-T7, SLT-T9 and SLT-T11
with those in the public databases as described above, revealed no significant homologies.
The sequences for SLT-T10 and SLT-T12 were found to show some homology to sequences
previously identified in humans.

20 The sequence of SLT-T1 was determined to show some homology to a PAC
clone of unknown protein function. The cDNA sequence of SLT-T1 (SEQ ID NO: 102) was
found to contain a mutator (MUT) domain. Such domains are known to function in removal
of damaged guanine from DNA that can cause A to G transversions (see, for example, el-
Deiry, W.S., 1997 *Curr. Opin. Oncol.* 9:79-87; Okamoto, K. et al. 1996 *Int. J. Cancer*
25 65:437-41; Wu, C. et al. 1995 *Biochem. Biophys. Res. Commun.* 214:1239-45; Porter, D.W.
et al. 1996 *Chem. Res. Toxicol.* 9:1375-81). SLT-T1 may thus be of use in the treatment, by
gene therapy, of lung cancers caused by, or associated with, a disruption in DNA repair.

Example 7

SYNTHESIS OF POLYPEPTIDES

Polypeptides may be synthesized on a Perkin Elmer/Applied Biosystems
5 Division 430A peptide synthesizer using FMOC chemistry with HPTU (O-Benzotriazole-
N,N,N',N'-tetramethyluronium hexafluorophosphate) activation. A Gly-Cys-Gly sequence
may be attached to the amino terminus of the peptide to provide a method of conjugation,
binding to an immobilized surface, or labeling of the peptide. Cleavage of the peptides from
the solid support may be carried out using the following cleavage mixture: trifluoroacetic
10 acid:ethanedithiol:thioanisole:water:phenol (40:1:2:2:3). After cleaving for 2 hours, the
peptides may be precipitated in cold methyl-t-butyl-ether. The peptide pellets may then be
dissolved in water containing 0.1% trifluoroacetic acid (TFA) and lyophilized prior to
purification by C18 reverse phase HPLC. A gradient of 0%-60% acetonitrile (containing
0.1% TFA) in water (containing 0.1% TFA) may be used to elute the peptides. Following
15 lyophilization of the pure fractions, the peptides may be characterized using electrospray or
other types of mass spectrometry and by amino acid analysis.

From the foregoing, it will be appreciated that, although specific embodiments
of the invention have been described herein for the purposes of illustration, various
20 modifications may be made without deviating from the spirit and scope of the invention.

9. A vaccine comprising the polypeptide of claim 2 and an immune response enhancer.

5 10. The vaccine of claim 9 wherein the immune response enhancer is an adjuvant.

11. A vaccine comprising the polynucleotide of claims 1 or 4 and an immune response enhancer.

10

12. The vaccine of claim 11 wherein the immune response enhancer is an adjuvant.

13. A pharmaceutical composition for the treatment of lung cancer
15 comprising a polypeptide and a physiologically acceptable carrier, the polypeptide comprising an immunogenic portion of a lung protein or of a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a sequence selected from the group consisting of:

(a) sequences recited in SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64,
20 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181;

(b) sequences complementary to the sequences of SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64, 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181; and

(c) variants of the sequences of (a) and (b).

25

14. A vaccine for the treatment of lung cancer comprising a polypeptide and an immune response enhancer, said polypeptide comprising an immunogenic portion of a lung protein or of a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a sequence selected from the group consisting of:

30 (a) sequences recited in SEQ ID NO: 12-18, 20, 21, 26, 49, 50, 52, 54, 64, 66, 68, 69, 71, 78, 84, 85, 88, 91, 92, 116-120, 126-138, 140-142, 150, 155 and 159-181;

21. A pharmaceutical composition comprising a fusion protein according to any one of claims 18-20 and a physiologically acceptable carrier.

5 22. A vaccine comprising a fusion protein according to any one of claims 18-20 and an immune response enhancer.

23. The vaccine of claim 22 wherein the immune response enhancer is an adjuvant.

10

24. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient an effective amount of the pharmaceutical composition of claim 21.

15 25. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient an effective amount of the vaccine of claim 22.

26. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient a polynucleotide under conditions such that the polynucleotide enters a cell of the patient and is expressed therein, the polynucleotide having a sequence selected from the group consisting of:

20

- (a) a sequence provided in SEQ ID NO: 102;
- (b) sequences complementary to a sequence of SEQ ID NO: 102; and
- (c) variants of the sequence of SEQ ID NO: 102.

25

27. A method for detecting lung cancer in a patient, comprising:

- (a) contacting a biological sample obtained from the patient with a binding agent which is capable of binding to a polypeptide, the polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, wherein said protein comprises an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences provided in SEQ ID NO: 1-31, 49-

30

- (a) sequences recited in SEQ ID NO: 1-11, 19, 22-25, 27-31, 51, 53, 55, 63, 70, 72, 79, 80, 86, 87, 89, 90, 102-107, 109, 139, 143-149, 151-154 and 156-158;
- (b) the complements of nucleotide sequences recited in SEQ ID NO: 1-11, 19, 22-25, 27-31, 51, 53, 55, 63, 70, 72, 79, 80, 86, 87, 89, 90, 102-107, 109, 139, 143-149, 151-154 and 156-158; and
- (c) variants of the sequences of (a) and (b).

32. A method for inhibiting the development of lung cancer in a patient, comprising administering to the patient a therapeutically effective amount of a monoclonal antibody according to claim 31.

33. The method of claim 32 wherein the monoclonal antibody is conjugated to a therapeutic agent.

34. A method for detecting lung cancer in a patient comprising:

- (a) obtaining a biological sample from the patient;
- (b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof; and
- (c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers, thereby detecting lung cancer.

35. The method of claim 34, wherein at least one of the oligonucleotide primers comprises at least about 10 contiguous nucleotides of a polynucleotide comprising a sequence selected from SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181.

provided in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof.

44. A method for detecting lung cancer in a patient, comprising:

(a) obtaining a biological sample from the patient;

5 (b) contacting the biological sample with an oligonucleotide probe specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-
10 120 and 126-181, the complements of said nucleotide sequences and variants thereof; and
(c) detecting in the sample a DNA sequence that hybridizes to the oligonucleotide probe, thereby detecting lung cancer in the patient.

45. The method of claim 44 wherein the oligonucleotide probe comprises at least about 15 contiguous nucleotides of a polynucleotide having a nucleotide sequence
15 selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said nucleotide sequences and variants thereof.

46. A diagnostic kit comprising an oligonucleotide probe specific for a polynucleotide encoding a polypeptide comprising an immunogenic portion of a lung tumor
20 protein or a variant thereof, said protein comprising an amino acid sequence encoded by a polynucleotide comprising a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55, 63, 64, 66, 68-72, 78-80, 84-92, 102-110, 116-120 and 126-181, the complements of said sequences and variants thereof.

47. The diagnostic kit of claim 46, wherein the oligonucleotide probe
25 comprises at least about 15 contiguous nucleotides of a polynucleotide having a nucleotide sequence selected from the group consisting of sequences recited in SEQ ID NO: 1-31, 49-55,

pharmaceutically acceptable carrier.

55. A composition for the treatment of lung cancer in a patient, comprising T cells proliferated in the presence of a polynucleotide of claim 1, in combination with a pharmaceutically acceptable carrier.

56. A method for treating lung cancer in a patient, comprising the steps of:
- (a) incubating antigen presenting cells in the presence of at least one polypeptide of claim 2; and
- 10 (b) administering to the patient the incubated antigen presenting cells.

57. A method for treating lung cancer in a patient, comprising the steps of:
- (a) incubating antigen presenting cells in the presence of at least one polynucleotide of claim 1; and
- 15 (b) administering to the patient the incubated antigen presenting cells.

58. The method of claims 54 or 55 wherein the antigen presenting cells are selected from the group consisting of dendritic cells and macrophage cells.

- 20 59. A composition for the treatment of lung cancer in a patient, comprising antigen presenting cells incubated in the presence of a polypeptide of claim 2, in combination with a pharmaceutically acceptable carrier.

60. A composition for the treatment of lung cancer in a patient, comprising
- 25 antigen presenting cells incubated in the presence of a polynucleotide of claim 1, in combination with a pharmaceutically acceptable carrier.

SEQUENCE LISTING

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taaaatnggt cctcctwaaa aaagttcggc tggggcccgc tggctcacat tttkttaycc 540
cycccccttt tggggarggc caarggccgg kttgawtnnc ccttgagggg ccanaactcc 600
agnaaccrgn cccgggccc arsmgwkstr armcccttcc cyyccmaraa aawwcsmaa 660
wwttyccsc cygsykgggt gggkascggt myyyyygntm csyagcttgc tt 712

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<210> 5

<211> 679

<212> DNA

<213> Homo sapiens

<400> 5

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cagaggggtc tgcaggatgt gctattttta agcagctggg tgcaacttgt gaaaacggga 180
atctngaagc agaacatgtn atcagcgatg gctgggattg gtggacagga ttgacaggag 240
tatttgaggc tctaccaggc ctgtctacag gacagcttca tcgaaggac attttttaac 300
ctgttatttt anatnccaca tatntttttt aatgctnaag catacagggt gaatttctgg 360
atcgtaacta ctagtactt ctgaggttta cagtngaag atgttctcnn aggtttatca 420
agttntgtta ttgatgatng gtaatctaca cctctggaag ctgtngaag tgaaaaagat 480
ncntncant gaccagtttg nagggcactc tcttctggna agnaatccgn ccaaaaaaat 540
tgtttchnagg gggcntgggg ggtttaaaaa aatgtttctn ttncntaaa aatgtttacc 600
cnnctattga aaaaatgggg gtcngggggg gcttnaaatc ccnnttntt gaattntnta 660
tccggaant tggtttccc 679

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<210> 6

<211> 369

<212> DNA

<213> Homo sapiens

<400> 6

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tcagtccagt catgggtcct ataagagaag tcaactctgtg agtttccatg gaggaagaaa 60
aagcttcatt tctttaccct gcagcaacag cggaggagg gagagcctat cttctttgca 120
aattcattaa ctttgtggtt gaagggagca gcgtcngaaa ctgcttttagc acagtgggag 180
gaaaacaaac agattcatct ccggaaacca aaggaaagg ttagtgggtt tttattagcc 240
agctgtatcc tagatggtca atttcacgtg gatgaatata ccttacgtac gtttctcttg 300
cttcctacct nggcctgatc agctnggcac ttraatcatt ccgtnggggt wgctgtnaca 360
ctggactga 369

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<210> 7

<211> 264

<212> DNA

<213> Homo sapiens

<400> 7

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tgctggatra gggatggggc acgggagcac agatmgactt taactgcccc cacgtntcm 60
aggaaaggat tacaggcgtg agccactgcg cccggcctct tctccacttt cataggttcc 120
agtctctggt tcttctttct cagtttggtt tttttgcttc ttaaammatg gagatnagaa 180
tgaacactac actcggatc aggaagccct gcctggcgcc tctgtcacct gtctaggggc 240
ttcttctcac tgagtcaccc agca 264

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<210> 8

<211> 280

<212> DNA

<213> Homo sapiens

<400> 8

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acctcaactg cccanaacan aactgttgta caagatttga ggatttaaca atatttcaca 60
tgaaatattt cagacctacg ngagggttta aagacnaatt aaatgagcac cngtgtgccc 120
accgcccna ttaagaatta gagcaagcag tgagggtgaag ccttgctcctt gcttttaaca 180
tagaaagtga tccaaattca ccaaacttga cttngggttt tgcagtgtgg cctcctgatt 240
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<210> 9

<211> 449

<212> DNA

<213> Homo sapiens

<400> 9

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tcgtcaactc caggatggct ttgaaaatna atggacacag atctctcctg ttttgatrat 60
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ggttccctgt ctcccatggg cgtcatttca tgtgtctctc tgccttcccc cagatattct 180
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ggtacgggag gaaacactgg gctnaccag atnctggact tgagtcttgc ctctgctgct 300
tgctgcacag ctctgtcat ggtgctaaac ctgtgacctg cctcacaggc ttagagcatg 360
cccgtagaag tactctnaac taaratgctt tccacaaatg agatggtttc atgaaaactt 420
caaataagagg gcctgggcaa aaaaaaaaaa 449

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<210> 10

<211> 538

<212> DNA

<213> Homo sapiens

<400> 10

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tttttttttt ttcccaaagg cctcaraaca ctagtcttct aattccaagc agaaagttac 60

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atccgcccggg atacatgccca cttgggtttga taaatcaaaa tacagcatcc ttcagatccc 120
tttgcctgagc aatacaatta tttgtatatg ttactttttt ttctgtttgg ctnaaaagatt 180
tgatatgagc tgaggaaaat gaagccttta ctgctatnag atctnatccc tttccaccac 240
ctttcagggg tnttggcact gcayatatc agaattcccc nnagtcgctn gtgataaaaa 300
tgtcttcaga gatggcagaa tatgtttcct ttggtacatg ttcattaaaa atatacacgt 360
gctcactact gtggatatgt atgtnttgac cgatnacaca ggctgattta ggggaagagat 420
aaaagcacac ttngaattta ttagcctttc accnagacta anattctgaa attaagaatg 480
tattccttgg tcaacaattt tcctcttctc ttagccctct tacattgtan tggactga 538

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<210> 11

<211> 543

<212> DNA

<213> Homo sapiens

<400> 11

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tttttttttt ttgcccacag ctgccatctt tgtgtgataa ggccaacctt ctatgggaat 60
caaccctcgc catcccagca aatccctctc ctcccttctc atgggagtgc cttgtattca 120
tcaggcatct gggacttgat gtgggtntgg gatttgaaat cagagcacct nggtctctst 180
caccattctn tcacttatta gctctnacct tgggtnaata cctgccttag tgtcntaggt 240
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gggtaaaacc acaggggtact ccgctcctcc naagaatgga gaattttttc tagaagccca 360
natntgcttg gaaggttggc caccnagagc cnnaatcttc ttttatttnc cactgaangc 420
ctaagaggna attctgaact catcccnna tgacctctcc cgaatmagaa tatctctggc 480
acttaccata ttttcttgcc ctcttccact tacnaaactc ctttatttct taacnggacg 540
aaa 543

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<210> 12

<211> 329

<212> DNA

<213> Homo sapiens

<400> 12

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cgatgacttg ggcagtgagt gggcctcctg ccagggtggca gggcacagct tagaccaaac 60
ccttggectc cccctctcgc agstacctct gaccaagaag gaaactagca agcctatgct 120
ggcaagacca taggtggggt gctgggaatc ctgggggccc gctggcaccc actcctgggtg 180
ctcaagggag agaccactt gttcagatgc atrggcctca ggcgggtcaa ggcrgtctta 240
gagccacaga gtcaaataaa aatcaatttt gagagaccac agcacctgct gctttgatcg 300
tgatgttcaa ggcaagttgc aagtcacgc 329

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<210> 13

<211> 314

<212> DNA

<213> Homo sapiens

<400> 13

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cgatgacttg caccggggag ctgtgacagt ggcctggaag cagatggcag ccccgtaag 60
gcgggagtgg agaccaccaa accctccaaa cagagcaaca actagtacgc ggccagcagc 120
tacctgagcc tgacgcccga gcagtggaa gttccacagaa gctacagctg ccagggtcacg 180
catgaaggga gcaccgtgga gaagacagt gcccctacag aatgttcata ggttcccnac 240
tctnacccca cccacgggag cctgganctg cangatccc ggggaagggg ctctctcccc 300
atcccaagtc atcg 314

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<210> 14

<211> 691

<212> DNA

<213> Homo sapiens

<400> 14

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cgattacttg cacaatgcan attagaaccc aaatgaaggg tacaaccag atcttctggc 60
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gaagagaaga ataaagtcta ttttggtctt tggtagcchg ggtaangaga atgctstcac 180
tctacnagaa aaccnnaagt gaaccgggt aatcaggacc gtgcttggga agggagcagg 240
ggcattacct ttcaacacca gaggttcttt gccttctctc tgcagggact cgargactat 300
gtgaagtggc tgggarggca tcactcggct tgggttcattg gtrttctcat cataaactat 360
natttctttg gaaaaagatc ctcttgaaag artccttgcc ttccctacag gaaatcaagt 420
ctaggacagt gatcttgccc ctgcttgcas tctccgccgg ctgatcttat csgsccaggt 480
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ccnwtnaatg amssggggccc ttaactccgg scrpgtnamy ncttgsctsc rattttgggt 600
ycytcttctt ttgscmagg tcktcnaaac cacttngttr aattccccgg sccgcctkgc 660
nggtycaacc wttttgggaa mamcycccc c 691

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<210> 15

<211> 355

<212> DNA

<213> Homo sapiens

<400> 15

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acctgaactg tgtgttgaag agtgatgtcc tgctgcctgg agctcaagtc actactgatg 60
accgtgccta tgtccgacag ctagttnect ccattggatgt gactgagacc aatgtcttct 120
tcyaccctcg gctcttacct ttgacnaagt ctcccggtga gactactacc gaaccaccag 180
cagttcgagc ctctnaagag cgtctaagcg atggggatat atatttactg gagaatgggc 240
tcaacctctt cctctgggtg ggagcaagcg tccagcaggg tgttgtccag agccttttca 300
gcgtctctc ctccagtcag atcaccagtg gtntgagtgt tctgccagtt caggt 355

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<210> 16

<211> 522

<212> DNA

<213> Homo sapiens

<400> 16

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tcagtcagtg gaggtggaag acttcgaggg tcgtggggagc cgcttctcca agtctgctga 60
tgagagacag cgcattgctg tgcagcgtan ggacgaactc ctccagcaag ctccgagacg 120
tttcttgaac aaaagttctg aagatgatgc ggctcagag agcttctctc cctcggaagg 180
tgctgctctt gaccccgta cctnccgtcg aangatgctg gctgccgccg cggaaacggan 240
gcttcagaag cagcagacct cctnccgtc ccttgccctc ctccagtgcc tccctgcgcc 300
tgtgcccggc tgactggagg aggcctgtcc aattctgccc gccccatgga aaagcgggct 360
tgactgcatt gccgctgtat naaagcatgt ggtcttacag tgttnggacn gctnatnaat 420
ttnatcctnc tntgtaatac ttctatgtg acatttctct tccccctgga aacactgcan 480
attttaactg tgagtttgat ctcttctnct gttactggac tg 522

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<210> 17

<211> 317

<212> DNA

<213> Homo sapiens

<400> 17

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gtgtcgcgaa ttccggtggt tgctaagaaa aggaagaaga agtcttacac cactcccaag 60
aaggataagc accagagaaa gaaggttcag ccggccgtcc tgaaatatta taagggtggat 120
gagaatggca aaattagttg ccttcgtcga gaggccccct ctgatgaatg tgggtgctggg 180
gtgtttatgg caagtcactt tgacagacat tattgtggca aatgttgtct gaccactgt 240

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ttcaactaac cagaagacaa gtaactgtat gagttaatta aagacatgaa ctaaaaaaaaa 300
 aaaaaaaaaa actcgag 317

<210> 18

<211> 392

<212> DNA

<213> Homo sapiens

<400> 18

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aggaacatgt	taaaaatcct	tacaaaggca	aaaaactcaa	gaaacaccca	gacttcccca	120
agaagcccct	gaccccttat	ttccgcttct	tcatggagaa	gcgggccaag	tatgcgaaac	180
tccaccctca	gatgagcaac	ctggacctga	ccaagattct	gtccaagaaa	tacaaggagc	240
ttccggagaa	gaagaagatg	aaatatgttc	cggacttcca	gagaagagaa	acaggagttc	300
gagcgaaacc	tggcccgatt	cagggaggat	cacccccacc	ttatccagaa	tgccaagaat	360
cggacatccc	agagaagccc	caagaccccc	cg			392

<210> 19

<211> 2624

<212> DNA

<213> Homo sapiens

<400> 19

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tcttgggctg	cccactgccc	gatcctaata	actattatca	ccgacgtaat	gagatgacca	180
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tcgatatcaa	caacaacttt	ccggatttaa	actcgtgct	ctgggaggca	gaggaccagc	660
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aataaaaatc	cactccagta	gtaactctgt	agcaggcttt	ccctgttgtt	ttgactgtaa	1620
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gtctgagatt	ctaaaaagg	tgcttgacca	ctggccagg	agggaaatca	ggccttcccc	1860
catttgcgtg	acattcaagc	ttcccagtc	atttgcaagt	ggcacagtgt	acattgcagc	1920

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accagggaa tcctttgccc cagatgttat catttgagat gctcttatgc agcctaagaa 1980
aatccatcct ctctggcccc aggggacaag ccaagctgct atgtacacac tcggtgttct 2040
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tgtgttgatc ctgctgcctt gttgacctgg agagaatgaa acaaacaac acataaacia 2580
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<210> 20

<211> 488

<212> DNA

<213> Homo sapiens

<400> 20

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ctttcaaccc gcgctcgccg gctccagccc cgcgcgcccc cacccttgc cctcccgccg 60
gctccgcagg gtgaggtggc ttgaccccg ggttgcccgg ccagcacgac cgaggaggtg 120
gctggacagc tggaggatga acggagaagc cgactgcccc acagacctgg aaatggccgc 180
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gaacaacctt ccatccaatg acagctccca gttcaaaaacc acccaaacac acatggaccg 300
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ctctaataag gtgaggaagt tccgtacatt gacagaattg atcctcgata ctcaggaaca 420
tgtttaaaat ccttaciaag gcaaaaaatc aagaaacacc ccgacttccc cgagaaagcc 480
cctaaccc 488

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<210> 21

<211> 391

<212> DNA

<213> Homo sapiens

<400> 21

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atggaattgt ggttttctct ttgggatcaa tgggtctcaga aattccagag aagaaagctg 60
tggcgattgc tgatgctttg ggcaaaatcc ctacagacgt cctgtggcgg tacactggaa 120
cccgaccatc gaactcttgc aacaacacga tacttgctca gtggctaccc caaacgata 180
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aaagcatatg caatggcgtt cccatgggtg tgataccctt atttgggtat cagatggaca 300
atgcaaagcg caggagact aaggagctg gagtgaacct gaatgttctg gagatgactt 360
ctgaagatct agaagatgct ctgaagagca g 391

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<210> 22

<211> 1320

<212> DNA

<213> Homo sapiens

<400> 22

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aatctgctgg gaatttcttg ggttgacagc tcttgatcc ctattttgaa cagtggtagt 60
gtcctggatt acttttcaga aagaagtaat cctttttatg acagaacatg taataatgaa 120
gtggtcaaaa tgcagaggct aacattagaa cacttgaatc agatggttgg aatcgagtac 180
atccttttgc atgctcaaga gccattctt tcatcattc ggaagcaaca gcggcagtc 240
cctgcccagg ttatcccact agctgattac tatatcattg ctggagtgat ctatcaggca 300
ccagacttgg gatcagttat aaactctaga gtgcttactg cagtgcattg tattcagtc 360

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gcttttgatg aagctatgtc atactgtcga tatcatcctt ccaaagggtg ttggtggcac 420
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agctctatct ttcagagaca acgtgtggat gctttacttt tagacctcag acaaaaattt 540
ccacccaaat ttgtgcagct aaagcctgga gaaaagcctg ttccagtggg tcaaacaaag 600
aaagaggcag aacctatacc agaaactgta aaacctgagg agaaggagac cacaagaat 660
gtacaacaga cagtgaagtgc taaaggcccc cctgaaaaac ggatgagact tcagtgaagta 720
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gtaacctttc ctttcccgga cttgagcaac ctacacactc acatgtttaa tggtagatat 1260
gttttaaagc aagataaagg tattggtttt aaaaaaaaaa aaaaaaaaaa aaaactcgag 1320

```

<210> 23

<211> 633

<212> DNA

<213> Homo sapiens

<400> 23

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tgatgggggt taagccgggg gaggcagcat cggggcctgc tgaagacctt gtgagaagat 120
ctgagaaaga tactgcagct gttgtctcca gacagggcag ctccctgaac ctcttgaaag 180
atgtgcagat cacagaacca gaagctgagc cagagtccaa gtctgaaccg agacctccaa 240
tttctctccc gagggctccc cagaccagag ctgtcaagcc ccgacttcat cctgtgaagc 300
caatgaatgc cacggccacc aaggttgcta actgcagctt gggaactgcc accatcatcg 360
gtgagaactt gaacaatgag gtcagtatga agaaatacag cccctcgagc cctgcatttg 420
catatgcgca gctgaccac gatgagctga ttgagctggt cctcaaacag aaggaaacga 480
taagcaagaa ggagttccag gtccgcgagc tgggaagacta cattgacaac ctgctcgtca 540
gggtcatgga agaaaccccc aatatcctcc gcaccccgac tcagggttggc aaaaaagcag 600
gaaagatgta aattagcaga aaaaaaactc gag 633

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<210> 24

<211> 1328

<212> DNA

<213> Homo sapiens

<400> 24

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gtaaacgctc tcggaattat ggcgccgggtg gatateccag acaatctgct gggaatttct 60
tggttgatg gctcttgat ccctattttg aacagtggta gtgtcctgga ttacttttca 120
gaaagaagta atccttttta tgacagaaca tgtaataatg aagtgggtcaa aatgcagagg 180
ctaacattag aacacttgaa tcagatgggt ggaatcgagt acatcctttt gcatgctcaa 240
gagccattc ttttcatcat tcggaagcaa cagcggcagt cccctgccc aagttatcca 300
ctagctgatt actatatcat tgctggagtg atctatcagg caccagactt gggatcagtt 360
ataaactcta gagtgtttac tgcagtgcac ggtattcagt cagcttttga tgaagctatg 420
tcatactgtc gatatcatcc ttccaaaggg tattggtggc acttcaaaga tcatgaagag 480
caagataaag tcagacctaa agccaaaagg aaagaagaac caagctctat ttttcagaga 540
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gtggatcaaa caaagaaaga ggcagaacct ataccagaaa ctgtaaaacc tgaggagaag 660
gagaccacaa agaattgtaca acagacagtg agtgctaaag gccccctga aaaacggatg 720
agacttcagt gagtactgga caaaagagaa gcctggaaga ctctcatgc tagttatcat 780
acctcagtac tgtggctctt gagctttgaa gtactttatt gtaaccttct tatttgtatg 840

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gaatgcgctt atttttttga aaggatatta ggccggatgt ggtggctcac gcctgtaatc 900
ccagcacttt gggaggccat ggcgggtgga tcacttgagg tcagaagttc aagaccagcc 960
tgaccaatat ggtgaaaccc cgtctctact aaaaatacaa aaattagccg ggcgtggtgg 1020
cgggcgcca tagtcccagc tactcgggag gctgagacag gagacttgct tgaacccggg 1080
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ttgtgatatg tcttctgtaa cctttcctct cccggacttg agcaacctac acactcacat 1260
gtttactggt agatatgttt aaaagcaaaa taaaggattt tgtataaaaa aaaaaaaaaa 1320
aaactcga                                     1328

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<210> 25

<211> 1758

<212> DNA

<213> Homo sapiens

<400> 25

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gttttttttt tttttttttt aaagagttgc aacaattcat ctttatttct tattttcctc 60
tggagatgca gaatttggtt tatttcaccc caagtatatt tgggatagtt ggctcctcgc 120
tgggtcagga tggctgggtg ccttctcccc tggcatggtt ctcttctctg cagggcgagg 180
ggcagggagc tagtaaaacc tcgcaatgac agccgcaatg gcagacccaa tggagcccag 240
gatgaacttg gtcaatccgg agagtccagt tgctcccagt gactgcagag tagccacaag 300
gctgcccgag gcaactccac cccatttggc aatggccgcc gcggacatca tcttggctgc 360
tatggaggag gaggcgattc ccgccgcagt gaagcccatg gcactgagtg gcggcggtgg 420
atatccgaga caatctgctg ggaatttctt ggggtgacag ctcttggatc cctattttga 480
acagtggtag tgtcctggat tacttttcag aaagaagtaa tcctttttat gacagaacat 540
gtaataatga agtggcctaa atgcagaggg taacattaga acacttgaat cagatgggtg 600
gaatcgagta catccttttg catgctcaag agccatttct tttcatcatt cggagccaac 660
agcggcagtc cctgcccac gttatcccac tagctgatta ctatatcatt gctggagtga 720
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gtattcagtc agcttttgat gaagctatgt catactgtcg atatcatcct tccaaagggt 840
attggtggca cttcaaagat catgaagagc aagataaagt cagacctaaa gccaaaagg 900
aagaagaacc aagctctatt tttcagagac aacgtgtgga tgctttactt ttagacctca 960
gacaaaaatt tccacccaaa tttgtgcagc taaagcctgg agaaaagcct gttccagtgg 1020
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tttgtctcaa aaaagaagaa aagatattat tcccatcatg atttcttgtg aatatttgtt 1620
atatgtcttc tgttaccttt cctctcccgg aattgagcaa cctacacact cacatgttta 1680
ctggtagata tgtttaaaag caaataaagg tattggtata tattgcttca aaaaaaaaaa 1740
aaaaaaaaa aactcgag                                     1758

```

<210> 26

<211> 493

<212> DNA

<213> Homo sapiens

<400> 26

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gagggcagcg gcagggcctg gtggcgagag cgcggtgtc actgcgccc agcatcccag 60
agctttccga gcggacgagc cggccgtgcc gggcatcccc agcctcgcta cctcgcagc 120

```

```

acacgtcgag ccccgccacag gcaaggggtcc ggaacttagc ccaaagcacg tttcccctgg 180
cagcgcgagga gacgcccggc cgcgcgcggg cgcacgcccc cctctcctcc tttgttccgg 240
gggtcgggcgg ccgctctcct gccagcgtcg ggatctcggc cccgggaggc gggccgtcgg 300
gcgcagccgc gaagattccg ttggaactga cgcagagccg agtgcagaag atctgggtgc 360
ccgtggacca caggccctcg ttgcccagat cctgtggggc aaagctgacc aactcccccg 420
ccgtcttcgt catggtgggc ctcccccgcc cggggcaaga cctacttctc cacgaaagct 480
tactcgctgc ctc

```

493

<210> 27

<211> 1331

<212> DNA

<213> Homo sapiens

<400> 27

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ggtggatc cgcagacaatc tgctgggaat ttcttgggtt gacagctctt ggatccctat 60
tttgaacagt ggtagtgtcc tggattactt ttcagaaaga agtaatcctt tttatgacag 120
aacatgtaat aatgaagtgg tcaaaatgca gaggctaaca ttagaacact tgaatcagat 180
ggttggaatc gagtacatcc ttttgcattc tcaagagccc attcttttca tcattcgga 240
gcaacagcgg cagtccccctg cccaagttat cccactagct gattactata tcattgctgg 300
agtgatctat caggcaccag acttggggtc agttataaac tctagagtgc ttactgcagt 360
gcatggattt cagtcagctt ttgatgaagc tatgtcctac tgcgatatc atccttccaa 420
agggatttgg tggcacttca aagatcatga agagcaagat aaagtcagac cttaaagccaa 480
aaggaaagaa gaaccaagct ctatttttca gagacaacgt gtggatgctt tactttttaga 540
cctcagaacaa aaatttccac ccaaatttgt gcagctaaag cctggagaaa agcctgttcc 600
agtggatcaa acaaagaaag aggcagaacc tataccagaa actgtaaaac ctgaggagaa 660
ggagaccaca aagaatgtac aacagacagt gagtgtctaa ggccccctg aaaaacggat 720
gagacttcag tgagtactgg acaaaagaga agcctggaag actcctcatg ctagtattca 780
tacctcagta ctgtggctct tgagctttga agtactttat tgtaaccttc ttatttgtat 840
ggaatgcgct tattttttga aaggatatta ggccggatgt ggtggctcac gcctgtaatc 900
ccagcacttt gggaggccat ggcgggtgga tcacttgagg tcagaagttc aagaccagcc 960
tgaccaatat ggtgaaaccc cgtctctact aaaaatacaa aaattagccg ggcgtggtgg 1020
cgggcgcccc tagtcccagc tactcgggag gctgagacag gagacttgct tgaaccggg 1080
aggtggaggt tgccttgagc tgattatcat gctgttgac tccagcttg ggcagagagc 1140
gagactttgt ctcaaaaaaa gaagaaaaga tattattccc atcatgattt cttgtgaata 1200
tttgttatat gtcttctgta acctttctc tcccggactt gagcaacctc cacactcaca 1260
tgtttactgg tagatatgtt taaaagcaaa ataaagggtat tggatataaa aaaaaaaaaa 1320
aaaaactcga g

```

1331

<210> 28

<211> 1333

<212> DNA

<213> Homo sapiens

<400> 28

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cggcgggtgga tatccgagac aatctgctgg gaatttcttg ggttgacagc tcttggatcc 60
ctattttgaa cagtggtagt gtcctggatt acttttcaga aagaagtaat cttttttatg 120
acagaacatg taataatgaa gtggtcaaaa tgcagaggct aacattagaa cacttgaatc 180
agatgggttg aatcagtagt atccttttgc atgctcaaga gccattctt ttcatcattc 240
ggaagcaaca gcggcagtc cctgccccag ttatccact agctgattac tatatcattg 300
ctggagtgat ctatcaggca ccagacttgg gatcagttat aaactctaga gtgcttactg 360
cagtgcattg tattcagtc gcttttgatg aagctatgtc atactgtcga tatcatcctt 420
ccaaagggta ttggtggcac ttcaaagatc atgaagagca agataaagtc agacctaaag 480
ccaaaggaa agaagaacca agctctattt ttcagagaca acgtgtggat gctttacttt 540
tagacctcag acaaaaattt ccacccaaat ttgtgcagct aaagcctgga gaaaagcctg 600
ttccagtgga tcaaacaaag aaagaggcag aacctatacc agaaactgta aaacctgagg 660

```

```

agaaggagac cacaaagaat gtacaacaga cagtgagtgc taaaggcccc cctgaaaaac 720
ggatgagact tcagtgagta ctggacaaaa gagaagcctg gaagactcct catgctagtt 780
atcatacctc agtactgtgg ctcttgagct ttgaagtact ttattgtaac cttcttattt 840
gtatggaatg cgcttatttt ttgaaaggat attaggccgg atgtgggtggc tcacgcctgt 900
aatcccagca ctttgggagg ccattggcggg tggatcactt gaggtcagaa gttcaagacc 960
agcctgacca atatggtgaa acccctgtctc tactaaaaat acaaaaatta gccgggctgt 1020
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cgggaggtgg aggttgccct gagctgatta tcatgctgtt gcactccagc ttgggcgaca 1140
gagcgagact ttgtctcaaa aaagaagaaa agatattatt cccatcatga tttcttgtga 1200
atatttgtga tatgtcttct gtaacctttc ctctcccgga cttgagcaac ctacacactc 1260
acatgtttac tggtagatat gtttaaaagc aaaataaagg tatttgtata aaaaaaaaaa 1320
aaaaaaaaac gag

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1333

<210> 29

<211> 813

<212> DNA

<213> Homo sapiens

<400> 29

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ctgagctgca cttcagcgaa ttcacctcgg ctgtggctga catgaagaac tccgtggcgg 60
accgagacaa cagccccagc tcctgtgctg gcctcttcat tgcttcacac atcgggtttg 120
actggccccg ggtctgggtc cacctggaca tcgctgctcc agtgcattgt ggcgagcgag 180
ccacaggctt tggggtggct ctctactgg ctctttttgg ccgtgcctcc gaggacccgc 240
tgctgaacct ggtatccccg ctggactgtg aggtggatgc ccaggaaagg gacaacatgg 300
ggcgtgactc caagagacgg aggtcgtgt gagggctact tcccagctgg tgacacaggg 360
ttccttacct cattttgcac tgactgattt taagcaattg aaagattaac taactcttaa 420
gatgagtttg gcttctcctt ctgtgcccag tggtgacagg agtgagccat tcttctctta 480
gaagcagctt aggggcttgg tggggtctgg agaaaattgt cacagacccc ataggtctcc 540
atctgtaagc tctgtccctt gtctccacc ctggctctta gagccacctc aggtcacctc 600
ctgtagttag tgtacttcct gacccaggcc cttgctcaag ctggggctcc ctggggtgtc 660
taaccagccc tgggtagatg tgactggctg ttagggaccc cattctgtga agcaggagac 720
cttcacagct cccaccaacc cccagttcac ttgaagttga attaaatat gccacaacat 780
aaaaaaaaaa aaaaaaaaaa aaaaaaacgc gag

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813

<210> 30

<211> 1316

<212> DNA

<213> Homo sapiens

<400> 30

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caggcgccca gtcattggccc aagagacagc accaccgtgt ggcccagtct caaggggtga 60
cagtcgaatc atagaaaaga tggaaaaaag gacatgtgcc ctgtgccctg aaggccacga 120
gtggagtcaa atatactttt caccatcagg aaatatagtt gctcatgaaa actgttttgt 180
gtattcatca ggactgggtg agtgtgagac tcttgatcta cgtaatacaa ttagaaactt 240
tgatgtcaaa tctgtaaaga aagagatctg gagaggaaga agattgaaat gctcattctg 300
taacaaagga ggcgccaccg tggggtgtga tttatggttc tgtaagaaga gttaccacta 360
tgtctgtgcc aaaaaggacc aagcaattct tcaagttgat ggaaaccatg gaacttacia 420
attattttgc ccagaacatt ctccagaaca agaagaggcc actgaaagtg ctgatgaccc 480
aagcatgaag aagaagagag gaaaaaacia acgctctca tcaggccctc ctgcacagcc 540
aaaaacgatg aaatgtagta acgccccaaag acatatgaca gaagagcctc atggtcacac 600
agatgcagct gtcaaatctc cttttcttaa ttcagttcat ggaagacttg tggatgagac 720
actatttgaa cacatactag aaaatatgga ttcagttcat ggaagacttg tggatgagac 780
tgcctcagag tcggactatg aagggtatga gaccttactg tttgactgtg gattatttaa 840
agacacacta agaaaattcc aagaagtaat caagagtaaa gcttgtgaat gggaagaaag 900
gcaaaggcag atgaagcagc agcttgaggc acttgacagc ttacaacaaa gcttgcgtcc

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atttcaagaa aatggggacc tggactgctc aagttctaca tcaggatcct tgctacctcc 960
tgaggaccac cagtaaaagc tgttcctcag gaaaactgga tggggcctcc atgttctcca 1020
aggatcgagg aagtcttcct gcctaccctg cccaccccag tcaagggcag caacaccaga 1080
gctttgctca gccttaaatg gaatcttaga gctttctctt gcttctgcta ctccctacaga 1140
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actgtgcatt gcacactgtt accatgggtt tatgtctact atcatatcac attgccataa 1260
tttagcacac ttaataaatg cttgtcaaaa cccaaaaaaa aaaaaaaaaa ctcgag 1316

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<210> 31

<211> 1355

<212> DNA

<213> Homo sapiens

<400> 31

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cggcgggtgga tatccgagac aatctgctgg gaatttcttg ggttgacagc tcttggatcc 60
ctatttttgaa cagtggtagt gtcctggatt acttttcaga aagaagtaat cctttttatg 120
acagaacatg taataatgaa gtgggtcaaaa tgcagaggct aacattagaa cacttgaatc 180
agatgggttg aatcgagtac atccttttgc atgctcaaga gccattctt ttcattctc 240
ggaagcaaca gcggcagtc cctgcccagg ttatcccact agctgattac tatatcattg 300
ctggagtgat ctatcaggca ccagacttgg gatcagttat aaactctaga gtgcttactg 360
cagtgcattg tattcagtca gcttttgatg aagctatgtc atactgtcga tatcatcctt 420
ccaaaagggtt ttggtggcac ttcaaagatc atgaagagca agataaagtc agacctaaag 480
ccaaaaggaa agaagaacca agctctatct ttcagagaca acgtgtggat gctttacttt 540
tagacctcag acaaaaatct ccacccaaat ttgtgcagct aaagcctgga gaaaagcctg 600
ttccagtggg tcaaacaag aaagaggcag aacctatacc agaaactgta aaacctgagg 660
agaaggagac cacaagaat gtacaacaga cagtgagtgc taaaggcccc cctgaaaaac 720
ggatgagact tcagttagta ctggacaaaa gagaagcctg gaagactcct catgctagtt 780
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cgggagggtg aggttgccct gagctgatta tcatgctgtt gcactccagc ttgggcgaca 1140
gaacgagact ttgtctcaaa aaaagaagaa aagatattat tcccatcatg atttcttgtg 1200
aatatttgtt atatgtcttc tggtaacctt tctctccccg gacttgaagc aacctcacac 1260
actcacatgt ttactggtag atatgtttta aaagcaaat aaaggatatt gtttttccaa 1320
aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa tcgag 1355

```

<210> 32

<211> 80

<212> PRT

<213> Homo sapiens

<400> 32

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Val Ser Arg Ile Arg Gly Gly Ala Lys Lys Arg Lys Lys Lys Ser Tyr
  1               5               10              15
Thr Thr Pro Lys Lys Asp Lys His Gln Arg Lys Lys Val Gln Pro Ala
 20              25              30
Val Leu Lys Tyr Tyr Lys Val Asp Glu Asn Gly Lys Ile Ser Cys Leu
 35              40              45
Arg Arg Glu Cys Pro Ser Asp Glu Cys Gly Ala Gly Val Phe Met Ala
 50              55              60

```

Ser His Phe Asp Arg His Tyr Cys Gly Lys Cys Cys Leu Thr His Cys
 65 70 75 80

<210> 33
 <211> 130
 <212> PRT
 <213> Homo sapiens

<400> 33
 Glu Ile Ser Asn Glu Val Arg Lys Phe Arg Thr Leu Thr Glu Leu Ile
 1 5 10 15
 Leu Asp Ala Gln Glu His Val Lys Asn Pro Tyr Lys Gly Lys Lys Leu
 20 25 30
 Lys Lys His Pro Asp Phe Pro Lys Lys Pro Leu Thr Pro Tyr Phe Arg
 35 40 45
 Phe Phe Met Glu Lys Arg Ala Lys Tyr Ala Lys Leu His Pro Gln Met
 50 55 60
 Ser Asn Leu Asp Leu Thr Lys Ile Leu Ser Lys Lys Tyr Lys Glu Leu
 65 70 75 80
 Pro Glu Lys Lys Lys Met Lys Tyr Val Pro Asp Phe Gln Arg Arg Glu
 85 90 95
 Thr Gly Val Arg Ala Lys Pro Gly Pro Ile Gln Gly Gly Ser Pro Pro
 100 105 110
 Pro Tyr Pro Glu Cys Gln Glu Ser Asp Ile Pro Glu Lys Pro Gln Asp
 115 120 125
 Pro Pro
 130

<210> 34
 <211> 506
 <212> PRT
 <213> Homo sapiens

<400> 34
 Asn Ser Glu Lys Glu Ile Pro Val Leu Asn Glu Leu Pro Val Pro Met
 1 5 10 15
 Val Ala Arg Tyr Ile Arg Ile Asn Pro Gln Ser Trp Phe Asp Asn Gly
 20 25 30
 Ser Ile Cys Met Arg Met Glu Ile Leu Gly Cys Pro Leu Pro Asp Pro

35	40	45
Asn Asn Tyr Tyr His Arg Arg Asn Glu Met Thr Thr Thr Asp Asp Leu		
50	55	60
Asp Phe Lys His His Asn Tyr Lys Glu Met Arg Gln Leu Met Lys Val		
65	70	75
Val Asn Glu Met Cys Pro Asn Ile Thr Arg Ile Tyr Asn Ile Gly Lys		
	85	90
Ser His Gln Gly Leu Lys Leu Tyr Ala Val Glu Ile Ser Asp His Pro		
	100	105
Gly Glu His Glu Val Gly Glu Pro Glu Phe His Tyr Ile Ala Gly Ala		
	115	120
His Gly Asn Glu Val Leu Gly Arg Glu Leu Leu Leu Leu Leu His		
	130	135
Phe Leu Cys Gln Glu Tyr Ser Ala Gln Asn Ala Arg Ile Val Arg Leu		
145	150	155
Val Glu Glu Thr Arg Ile His Ile Leu Pro Ser Leu Asn Pro Asp Gly		
	165	170
Tyr Glu Lys Ala Tyr Glu Gly Gly Ser Glu Leu Gly Gly Trp Ser Leu		
	180	185
Gly Arg Trp Thr His Asp Gly Ile Asp Ile Asn Asn Asn Phe Pro Asp		
	195	200
Leu Asn Ser Leu Leu Trp Glu Ala Glu Asp Gln Gln Asn Ala Pro Arg		
	210	215
Lys Val Pro Asn His Tyr Ile Ala Ile Pro Glu Trp Phe Leu Ser Glu		
225	230	235
Asn Ala Thr Val Ala Thr Glu Thr Arg Ala Val Ile Ala Trp Met Glu		
	245	250
Lys Ile Pro Phe Val Leu Gly Gly Asn Leu Gln Gly Gly Glu Leu Val		
	260	265
Val Ala Tyr Pro Tyr Asp Met Val Arg Ser Leu Trp Lys Thr Gln Glu		
	275	280
His Thr Pro Thr Pro Asp Asp His Val Phe Arg Trp Leu Ala Tyr Ser		
	290	295
Tyr Ala Ser Thr His Arg Leu Met Thr Asp Ala Arg Arg Arg Val Cys		
305	310	315
His Thr Glu Asp Phe Gln Lys Glu Glu Gly Thr Val Asn Gly Ala Ser		
	325	330
		335

Trp His Thr Val Ala Gly Ser Leu Asn Asp Phe Ser Tyr Leu His Thr
 340 345 350
 Asn Cys Phe Glu Leu Ser Ile Tyr Val Gly Cys Asp Lys Tyr Pro His
 355 360 365
 Glu Ser Glu Leu Pro Glu Glu Trp Glu Asn Asn Arg Glu Ser Leu Ile
 370 375 380
 Val Phe Met Glu Gln Val His Arg Gly Ile Lys Gly Ile Val Arg Asp
 385 390 395 400
 Leu Gln Gly Lys Gly Ile Ser Asn Ala Val Ile Ser Val Glu Gly Val
 405 410 415
 Asn His Asp Ile Arg Thr Ala Ser Asp Gly Asp Tyr Trp Arg Leu Leu
 420 425 430
 Asn Pro Gly Glu Tyr Val Val Thr Ala Lys Ala Glu Gly Phe Ile Thr
 435 440 445
 Ser Thr Lys Asn Cys Met Val Gly Tyr Asp Met Gly Ala Thr Arg Cys
 450 455 460
 Asp Phe Thr Leu Thr Lys Thr Asn Leu Ala Arg Ile Arg Glu Ile Met
 465 470 475 480
 Glu Thr Phe Gly Lys Gln Pro Val Ser Leu Pro Ser Arg Arg Leu Lys
 485 490 495
 Leu Arg Gly Arg Lys Arg Arg Gln Arg Gly
 500 505

<210> 35

<211> 96

<212> PRT

<213> Homo sapiens

<400> 35

Met Asn Gly Glu Ala Asp Cys Pro Thr Asp Leu Glu Met Ala Ala Pro
 1 5 10 15
 Arg Gly Gln Asp Arg Trp Ser Gln Glu Asp Met Leu Thr Leu Leu Glu
 20 25 30
 Cys Met Lys Asn Asn Leu Pro Ser Asn Asp Ser Ser Gln Phe Lys Thr
 35 40 45
 Thr Gln Thr His Met Asp Arg Glu Lys Val Ala Leu Lys Asp Phe Ser
 50 55 60
 Gly Asp Met Cys Lys Leu Lys Trp Val Glu Ile Ser Asn Glu Val Arg
 65 70 75 80

Lys Phe Arg Thr Leu Thr Glu Leu Ile Leu Asp Thr Gln Glu His Val
 85 90 95

<210> 36

<211> 129

<212> PRT

<213> Homo sapiens

<400> 36

Gly Ile Val Val Phe Ser Leu Gly Ser Met Val Ser Glu Ile Pro Glu
 1 5 10 15

Lys Lys Ala Val Ala Ile Ala Asp Ala Leu Gly Lys Ile Pro Gln Thr
 20 25 30

Val Leu Trp Arg Tyr Thr Gly Thr Arg Pro Ser Asn Leu Ala Asn Asn
 35 40 45

Thr Ile Leu Val Gln Trp Leu Pro Gln Asn Asp Leu Leu Gly His Pro
 50 55 60

Met Thr Arg Ala Phe Ile Thr His Ala Ser Ser His Gly Val Asn Glu
 65 70 75 80

Ser Ile Cys Asn Gly Val Pro Met Val Met Ile Pro Leu Phe Gly Asp
 85 90 95

Gln Met Asp Asn Ala Lys Arg Arg Glu Thr Lys Gly Ala Gly Val Thr
 100 105 110

Leu Asn Val Leu Glu Met Thr Ser Glu Asp Leu Glu Asp Ala Leu Lys
 115 120 125

Ser

<210> 37

<211> 238

<212> PRT

<213> Homo sapiens

<400> 37

Asn Leu Leu Gly Ile Ser Trp Val Asp Ser Ser Trp Ile Pro Ile Leu
 1 5 10 15

Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser Glu Arg Ser Asn Pro Phe
 20 25 30

Tyr Asp Arg Thr Cys Asn Asn Glu Val Val Lys Met Gln Arg Leu Thr
 35 40 45

Leu Glu His Leu Asn Gln Met Val Gly Ile Glu Tyr Ile Leu Leu His
 50 55 60

Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg Lys Gln Gln Arg Gln Ser
 65 70 75 80
 Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr Tyr Ile Ile Ala Gly Val
 85 90 95
 Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val Ile Asn Ser Arg Val Leu
 100 105 110
 Thr Ala Val His Gly Ile Gln Ser Ala Phe Asp Glu Ala Met Ser Tyr
 115 120 125
 Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp Trp His Phe Lys Asp His
 130 135 140
 Glu Glu Gln Asp Lys Val Arg Pro Lys Ala Lys Arg Lys Glu Glu Pro
 145 150 155 160
 Ser Ser Ile Phe Gln Arg Gln Arg Val Asp Ala Leu Leu Leu Asp Leu
 165 170 175
 Arg Gln Lys Phe Pro Pro Lys Phe Val Gln Leu Lys Pro Gly Glu Lys
 180 185 190
 Pro Val Pro Val Asp Gln Thr Lys Lys Glu Ala Glu Pro Ile Pro Glu
 195 200 205
 Thr Val Lys Pro Glu Glu Lys Glu Thr Thr Lys Asn Val Gln Gln Thr
 210 215 220
 Val Ser Ala Lys Gly Pro Pro Glu Lys Arg Met Arg Leu Gln
 225 230 235

 <210> 38
 <211> 202
 <212> PRT
 <213> Homo sapiens

 <400> 38
 Lys Gly Ser Glu Gly Glu Asn Pro Leu Thr Val Pro Gly Arg Glu Lys
 1 5 10 15
 Glu Gly Met Leu Met Gly Val Lys Pro Gly Glu Asp Ala Ser Gly Pro
 20 25 30
 Ala Glu Asp Leu Val Arg Arg Ser Glu Lys Asp Thr Ala Ala Val Val
 35 40 45
 Ser Arg Gln Gly Ser Ser Leu Asn Leu Phe Glu Asp Val Gln Ile Thr
 50 55 60
 Glu Pro Glu Ala Glu Pro Glu Ser Lys Ser Glu Pro Arg Pro Pro Ile
 65 70 75 80

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<210> 39
<211> 243
<212> PRT
<213> Homo sapiens
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<400> 39
Val Asn Ala Leu Gly Ile Met Ala Ala Val Asp Ile Arg Asp Asn Leu
  1              5              10              15
Leu Gly Ile Ser Trp Val Asp Ser Ser Trp Ile Pro Ile Leu Asn Ser
              20              25              30
Gly Ser Val Leu Asp Tyr Phe Ser Glu Arg Ser Asn Pro Phe Tyr Asp
              35              40              45
Arg Thr Cys Asn Asn Glu Val Val Lys Met Gln Arg Leu Thr Leu Glu
  50              55              60
His Leu Asn Gln Met Val Gly Ile Glu Tyr Ile Leu Leu His Ala Gln
  65              70              75              80
Glu Pro Ile Leu Phe Ile Ile Arg Lys Gln Gln Arg Gln Ser Pro Ala
              85              90              95
Gln Val Ile Pro Leu Ala Asp Tyr Tyr Ile Ile Ala Gly Val Ile Tyr
              100              105              110
Gln Ala Pro Asp Leu Gly Ser Val Ile Asn Ser Arg Val Leu Thr Ala
              115              120              125

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Val His Gly Ile Gln Ser Ala Phe Asp Glu Ala Met Ser Tyr Cys Arg
 130 135 140

Tyr His Pro Ser Lys Gly Tyr Trp Trp His Phe Lys Asp His Glu Glu
 145 150 155 160

Gln Asp Lys Val Arg Pro Lys Ala Lys Arg Lys Glu Glu Pro Ser Ser
 165 170 175

Ile Phe Gln Arg Gln Arg Val Asp Ala Leu Leu Leu Asp Leu Arg Gln
 180 185 190

Lys Ile Ser Thr Gln Ile Cys Ala Val Asp Gln Thr Lys Lys Glu Ala
 195 200 205

Glu Pro Ile Pro Glu Thr Val Lys Pro Glu Glu Lys Glu Thr Thr Lys
 210 215 220

Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys Arg Met
 225 230 235 240

Arg Leu Gln

<210> 40
 <211> 245
 <212> PRT
 <213> Homo sapiens

<400> 40
 Ala Ala Val Asp Ile Arg Asp Asn Leu Leu Gly Ile Ser Trp Val Asp
 1 5 10 15

Ser Ser Trp Ile Pro Ile Leu Asn Ser Gly Ser Val Leu Asp Tyr Phe
 20 25 30

Ser Glu Arg Ser Asn Pro Phe Tyr Asp Arg Thr Cys Asn Asn Glu Val
 35 40 45

Val Lys Met Gln Arg Leu Thr Leu Glu His Leu Asn Gln Met Val Gly
 50 55 60

Ile Glu Tyr Ile Leu Leu His Ala Gln Glu Pro Ile Leu Phe Ile Ile
 65 70 75 80

Arg Lys Gln Gln Arg Gln Ser Pro Ala Gln Val Ile Pro Leu Ala Asp
 85 90 95

Tyr Tyr Ile Ile Ala Gly Val Ile Tyr Gln Ala Pro Asp Leu Gly Ser
 100 105 110

Val Ile Asn Ser Arg Val Leu Thr Ala Val His Gly Ile Gln Ser Ala
 115 120 125

Phe Asp Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr
 130 135 140
 Trp Trp His Phe Lys Asp His Glu Glu Gln Asp Lys Val Arg Pro Lys
 145 150 155 160
 Ala Lys Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val
 165 170 175
 Asp Ala Leu Leu Leu Asp Leu Arg Gln Lys Phe Pro Pro Lys Phe Val
 180 185 190
 Gln Leu Lys Pro Gly Glu Lys Pro Val Pro Val Asp Gln Thr Lys Lys
 195 200 205
 Glu Ala Glu Pro Ile Pro Glu Thr Val Lys Pro Glu Glu Lys Glu Thr
 210 215 220
 Thr Lys Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys
 225 230 235 240
 Arg Met Arg Leu Gln
 245
 <210> 41
 <211> 163
 <212> PRT
 <213> Homo sapiens
 <400> 41
 Gly Glu Arg Gln Gly Leu Val Ala Arg Ala Arg Leu Ser Leu Arg Pro
 1 5 10 15
 Ser Ile Pro Glu Leu Ser Glu Arg Thr Ser Arg Pro Cys Arg Ala Ser
 20 25 30
 Pro Ala Ser Leu Pro Ser Gln His Thr Ser Ser Pro Ala Gln Ala Arg
 35 40 45
 Val Arg Asn Leu Ala Gln Ser Thr Phe Pro Leu Ala Ala Gln Glu Thr
 50 55 60
 Pro Gly Arg Ala Pro Ala His Ala Pro Leu Ser Ser Phe Val Pro Gly
 65 70 75 80
 Val Gly Gly Arg Ser Pro Ala Ser Val Gly Ile Ser Ala Pro Gly Gly
 85 90 95
 Gly Pro Ser Gly Ala Ala Ala Lys Ile Pro Leu Glu Leu Thr Gln Ser
 100 105 110
 Arg Val Gln Lys Ile Trp Val Pro Val Asp His Arg Pro Ser Leu Pro
 115 120 125
 Arg Ser Cys Gly Pro Lys Leu Thr Asn Ser Pro Ala Val Phe Val Met

130 135 140
 Val Gly Leu Pro Arg Pro Gly Gln Asp Leu Leu Leu His Glu Ser Leu
 145 150 155 160
 Leu Ala Ala

 <210> 42
 <211> 243
 <212> PRT
 <213> Homo sapiens

 <400> 42
 Val Asp Ile Arg Asp Asn Leu Leu Gly Ile Ser Trp Val Asp Ser Ser
 1 5 10 15
 Trp Ile Pro Ile Leu Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser Glu
 20 25 30
 Arg Ser Asn Pro Phe Tyr Asp Arg Thr Cys Asn Asn Glu Val Val Lys
 35 40 45
 Met Gln Arg Leu Thr Leu Glu His Leu Asn Gln Met Val Gly Ile Glu
 50 55 60
 Tyr Ile Leu Leu His Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg Lys
 65 70 75 80
 Gln Gln Arg Gln Ser Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr Tyr
 85 90 95
 Ile Ile Ala Gly Val Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val Ile
 100 105 110
 Asn Ser Arg Val Leu Thr Ala Val His Gly Ile Gln Ser Ala Phe Asp
 115 120 125
 Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp Trp
 130 135 140
 His Phe Lys Asp His Glu Glu Gln Asp Lys Val Arg Pro Lys Ala Lys
 145 150 155 160
 Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val Asp Ala
 165 170 175
 Leu Leu Leu Asp Leu Arg Gln Lys Phe Pro Pro Lys Phe Val Gln Leu
 180 185 190
 Lys Pro Gly Glu Lys Pro Val Pro Val Asp Gln Thr Lys Lys Glu Ala
 195 200 205
 Glu Pro Ile Pro Glu Thr Val Lys Pro Glu Glu Lys Glu Thr Thr Lys
 210 215 220

Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys Arg Met
 225 230 235 240

Arg Leu Gln

<210> 43

<211> 244

<212> PRT

<213> Homo sapiens

<400> 43

Ala Val Asp Ile Arg Asp Asn Leu Leu Gly Ile Ser Trp Val Asp Ser
 1 5 10 15

Ser Trp Ile Pro Ile Leu Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser
 20 25 30

Glu Arg Ser Asn Pro Phe Tyr Asp Arg Thr Cys Asn Asn Glu Val Val
 35 40 45

Lys Met Gln Arg Leu Thr Leu Glu His Leu Asn Gln Met Val Gly Ile
 50 55 60

Glu Tyr Ile Leu Leu His Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg
 65 70 75 80

Lys Gln Gln Arg Gln Ser Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr
 85 90 95

Tyr Ile Ile Ala Gly Val Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val
 100 105 110

Ile Asn Ser Arg Val Leu Thr Ala Val His Gly Ile Gln Ser Ala Phe
 115 120 125

Asp Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp
 130 135 140

Trp His Phe Lys Asp His Glu Glu Gln Asp Lys Val Arg Pro Lys Ala
 145 150 155 160

Lys Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val Asp
 165 170 175

Ala Leu Leu Leu Asp Leu Arg Gln Lys Phe Pro Pro Lys Phe Val Gln
 180 185 190

Leu Lys Pro Gly Glu Lys Pro Val Pro Val Asp Gln Thr Lys Lys Glu
 195 200 205

Ala Glu Pro Ile Pro Glu Thr Val Lys Pro Glu Glu Lys Glu Thr Thr
 210 215 220

Lys Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys Arg
 225 230 235 240

Met Arg Leu Gln

<210> 44

<211> 109

<212> PRT

<213> Homo sapiens

<400> 44

Glu Leu His Phe Ser Glu Phe Thr Ser Ala Val Ala Asp Met Lys Asn
 1 5 10 15

Ser Val Ala Asp Arg Asp Asn Ser Pro Ser Ser Cys Ala Gly Leu Phe
 20 25 30

Ile Ala Ser His Ile Gly Phe Asp Trp Pro Gly Val Trp Val His Leu
 35 40 45

Asp Ile Ala Ala Pro Val His Ala Gly Glu Arg Ala Thr Gly Phe Gly
 50 55 60

Val Ala Leu Leu Leu Ala Leu Phe Gly Arg Ala Ser Glu Asp Pro Leu
 65 70 75 80

Leu Asn Leu Val Ser Pro Leu Asp Cys Glu Val Asp Ala Gln Glu Gly
 85 90 95

Asp Asn Met Gly Arg Asp Ser Lys Arg Arg Arg Leu Val
 100 105

<210> 45

<211> 324

<212> PRT

<213> Homo sapiens

<400> 45

Arg Arg Pro Val Met Ala Gln Glu Thr Ala Pro Pro Cys Gly Pro Val
 1 5 10 15

Ser Arg Gly Asp Ser Pro Ile Ile Glu Lys Met Glu Lys Arg Thr Cys
 20 25 30

Ala Leu Cys Pro Glu Gly His Glu Trp Ser Gln Ile Tyr Phe Ser Pro
 35 40 45

Ser Gly Asn Ile Val Ala His Glu Asn Cys Leu Leu Tyr Ser Ser Gly
 50 55 60

Leu Val Glu Cys Glu Thr Leu Asp Leu Arg Asn Thr Ile Arg Asn Phe
 65 70 75 80

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<210> 46
<211> 244
<212> PRT
<213> Homo sapiens
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<400> 46
Ala Val Asp Ile Arg Asp Asn Leu Leu Gly Ile Ser Trp Val Asp Ser

25

1	5	10	15
Ser Trp Ile Pro Ile Leu Asn Ser Gly Ser Val Leu Asp Tyr Phe Ser	20	25	30
Glu Arg Ser Asn Pro Phe Tyr Asp Arg Thr Cys Asn Asn Glu Val Val	35	40	45
Lys Met Gln Arg Leu Thr Leu Glu His Leu Asn Gln Met Val Gly Ile	50	55	60
Glu Tyr Ile Leu Leu His Ala Gln Glu Pro Ile Leu Phe Ile Ile Arg	65	70	75
Lys Gln Gln Arg Gln Ser Pro Ala Gln Val Ile Pro Leu Ala Asp Tyr	85	90	95
Tyr Ile Ile Ala Gly Val Ile Tyr Gln Ala Pro Asp Leu Gly Ser Val	100	105	110
Ile Asn Ser Arg Val Leu Thr Ala Val His Gly Ile Gln Ser Ala Phe	115	120	125
Asp Glu Ala Met Ser Tyr Cys Arg Tyr His Pro Ser Lys Gly Tyr Trp	130	135	140
Trp His Phe Lys Asp His Glu Glu Gln Asp Lys Val Arg Pro Lys Ala	145	150	155
Lys Arg Lys Glu Glu Pro Ser Ser Ile Phe Gln Arg Gln Arg Val Asp	165	170	175
Ala Leu Leu Leu Asp Leu Arg Gln Lys Phe Pro Pro Lys Phe Val Gln	180	185	190
Leu Lys Pro Gly Glu Lys Pro Val Pro Val Asp Gln Thr Lys Lys Glu	195	200	205
Ala Glu Pro Ile Pro Glu Thr Val Lys Pro Glu Glu Lys Glu Thr Thr	210	215	220
Lys Asn Val Gln Gln Thr Val Ser Ala Lys Gly Pro Pro Glu Lys Arg	225	230	235
Met Arg Leu Gln			240

<210> 47

<211> 14

<212> DNA

<213> Homo sapiens

<400> 47

tttttttttt ttag

<210> 48
 <211> 10
 <212> DNA
 <213> Homo sapiens

<400> 48
 cttcaacctc

10

<210> 49
 <211> 496
 <212> DNA
 <213> Homo sapiens

<400> 49
 gcaccatgta ccgagcactt cggctcctcg cgcgctcgcg tccctcgtg cgggctccag 60
 ccgcagcctt agcttcggct cccggcttgg gtggcgcggc cgtgccctcg ttttggcctc 120
 cgaacgcggc tcgaatggca agccaaaatt ccttcggat agaatatgat acctttggtg 180
 aactaaaggt gccaaatgat aagtattatg gcgcccagac cgtgagatct acgatgaact 240
 ttaagattgg aggtgtgaca gaacgcgatgc caacccagc tattaagct tttggcatct 300
 tgaagcgagc ggccgctgaa gtaaacccag attatggctc tgatccaaag attgctaatt 360
 caataatgaa ggcagcagat gaggtagctg aaggtaaatt aaatgatcat tttcctctcg 420
 tggatggca gactggatca ggaactcaga caaatatgaa tgtaaataaa gtcattagcc 480
 aatagagcaa ttgaaa 496

<210> 50
 <211> 499
 <212> DNA
 <213> Homo sapiens

<400> 50
 agaaaaagtc tatgtttgca gaaatacaga tccaagacaa agacaggatg ggcactgctg 60
 gaaaagtatt taaatgcaaa gcagctgtgc tttgggagca gaagcaacc tttccattg 120
 aggaaataga agttgcccc acaaagacta aagaagttcg cattaagatt ttggccacag 180
 gaatctgtcg cacagatgac catgtgataa aaggaacaat ggtgtccaag tttccagtga 240
 ttgtgggaca tgaggcaact gggattgtag agagcattgg agaaggagt actacagtga 300
 aaccaggtga caaagtcac cctctctttc tgccacaatg tagagaatgc aatgcttgct 360
 gcaaccacaga tggcaacctt tgcattagga gcgatattac tggctcgtga gtactggctg 420
 atggcaccac cagatttaca tgcaagggcg aaccagtcca ccacttcattg aacaccagta 480
 catttaccga gtacacagt 499

<210> 51
 <211> 887
 <212> DNA
 <213> Homo sapiens

<400> 51
 gagtctgagc agaaaggaaa agcagccttg gcagccacgt tagaggaata caaagccaca 60
 gtggccagt accagataga gatgaatcgc ctgaaggctc agctggagaa tgaaaagcag 120
 aaagtggcag agctgtattc tatccataac tctggagaca aatctgatat tcaggacctc 180
 ctggagagtgc tcaggctgga caaagaaaaa gcagagactt tggctagtag cttgcaggaa 240
 gatctggctc ataccgaaa tgatgccaat cgattacagg atgccattgc taaggtagag 300
 gatgaatacc gagccttcca agaagaagct aagaaacaaa ttgaagattt gaatatgacg 360
 ttagaaaaat taagatcaga cctggatgaa aaagaaacag aaaggagtga catgaaagaa 420
 accatctttg aacttgaaga tgaagtagaa caacatcgtg ctgtgaaact tcatgacaac 480
 ctcatatttt ctgatctaga gaatacagtt aaaaaactcc aggaccaaaa gcacgacatg 540

gaaagagaaa taaagacact ccacagaaga cttcggaag aatctgcgga atggcggcag 600
 tttcaggctg atctccagac tgcagtagtc attgcaaag acattaaatc tgaagcccaa 660
 gaggagattg gtgatctaaa gcgccggtta catgaggctc aagaaaaaaa tgagaaactc 720
 acaaaagaat tggaggaaat aaagtcacgc aagcaagagg aggagcgagg cgggtatata 780
 attacatgaa tgccgttgag agagatttgg cagccttaag gcagggaatg ggactgagta 840
 gaaggtcctc gacttcctca gagccaactc ctacagtaaa aaccctc 887

<210> 52

<211> 491

<212> DNA

<213> Homo sapiens

<400> 52

ggcacgagct tttccaaaaa tcatgtctgt cttttctcta aagttcttac attttataga 60
 aaggaacctt tcaactcttga ggcctactac agctctcctc aggatttgcc ctatccagat 120
 cctgctatag ctcaagtttc agttcagaaa gtcactcctc agtctgatgg ctccagttca 180
 aaagtgaag tcaaagttcg agtaaatgtc catggcattt tcagtgtgtc cagtgcattc 240
 ttagtggagg ttcacaagtc tgaggaaaat gaggagccaa tggaaacaga tcagaatgca 300
 aaggaggaag agaagatgca agtggaccag gaggaaccac atgttgaaga gcaacagcag 360
 cagacaccag gcagaaaata aggcagagtc tgaagaaatg gagacctctc aagctggatc 420
 caaggataaa aagatggacc aaccacccca agccaagaag gcaaaagtga agaccagtac 480
 tgtggacctg g 491

<210> 53

<211> 787

<212> DNA

<213> Homo sapiens

<400> 53

aagcagttga gtaggcagaa aaaagaacct cttcattaag gattaaaatg tataggccag 60
 cacgtgtaac ttcgacttca agatttctga atccatagt agtatgtttc attgtcgtcg 120
 caggggtagt gatcctggca gtcaccatag ctctacttgt ttacttttta gcttttgatc 180
 aaaaatctta cttttatagg agcagttttc aactcctaaa tgttgaatat aatagtcagt 240
 taaattcacc agctacacag gaatacagga ctttgagtgg aagaattgaa tctctgatta 300
 ctaaaacatt caaagaatca aatttaagaa atcagttcat cagagctcat gttgccaaac 360
 tgaggcaaga tggtagtggt gtgagagcgg atgttgtcat gaaatttcaa ttcactagaa 420
 ataacaatgg agcatcaatg aaaagcagaa ttgagtctgt tttacgacaa atgctgaata 480
 actctggaaa cctggaaata aacccttcaa ctgagataac atcaattact gaccaggctg 540
 cagcaaattg gcttattaat gaatgtgggg ccggtccaga cctaataaca ttgtctgagc 600
 agagaatcct tggaggcact gaggtgagg aggggaagctg gccgtggcaa gtcagtctgc 660
 ggctcaataa tgcccaccac tgtggaggca gcctgatcaa taacatgtgg atcctgacag 720
 cagctcactg cttcagaagc aactctaate ctcgtgactg gattgccacg tctggtattt 780
 ccacaac 787

<210> 54

<211> 386

<212> DNA

<213> Homo sapiens

<400> 54

ggcattttca gtgtgtccag tgcattctta gtggagggtc acaagtctga ggaaaatgag 60
 gagccaatgg aaacagatca gaatgcaaag gaggaagaga agatgcaagt ggaccaggag 120
 gaaccacatg ttgaagagca acagcagcag acaccagcag aaaataaggc agagtctgaa 180
 gaaatggaga cctctcaagc tggatccaag gataaaaaga tggaccaacc accccaagcc 240
 aagaaggcaa aagtgaagac cagtactgtg gacctgccaa tcgagaatca gctattatgg 300

cagatagaca gagagatgct caacttgtag attgaaaatg agggtaagat gatcatgcag 360
gataaactgg agaaggagcg gaatga 386

<210> 55

<211> 1462

<212> DNA

<213> Homo sapiens

<400> 55

aagcagttga gtaggcagaa aaaagaacct cttcattaag gattaaaatg tataggccag 60
cacgtgtaac ttcgacttca agatttctga atccatatgt agtatgtttc attgtcgtcg 120
caggggtagt gatcctggca gtcaccatag ctctacttgt ttacttttta gcttttgatc 180
aaaaatctta cttttatagg agcagttttc aactcctaaa tgttgaatat aatagtcagt 240
taaattcacc agctacacag gaatacagga ctttgagtgg aagaattgaa tctctgatta 300
ctaaaacatt caaagaatca aatttaagaa atcagttcat cagagctcat gttgccaaac 360
tgaggcaaga tggtagtggt gtgagagcgg atgttgatcat gaaatttcaa ttcactagaa 420
ataacaatgg agcatcaatg aaaagcagaa ttgagtctgt tttacgacaa atgctgaata 480
actctggaaa cctggaaata aacccttcaa ctgagataac atcacttact gaccaggtcg 540
cagcaaatgg gcttattaat gaatgtgggg ccggtccaga cctaataaca ttgtctgagc 600
agagaatcct tggaggcact gaggctgagg agggaaagctg gccgtggcaa gtcagtctgc 660
ggctcaataa tgcccaccac tgtggaggca gcctgatcaa taacatgtgg atcctgacag 720
cagctcactg cttcagaagc aactctaate ctgctgactg gattgccacg tctgggtattt 780
ccacaacatt tcctaaacta agaagagag taagaaatat ttttaattcat aacaattata 840
aatctgcaac tcatgaaaat gacattgcac ttgtgagact tgagaacagt gtcaccttta 900
ccaaagatat ccatagtgtg tgtctcccag ctgctaccca gaattattcca cctgggtcta 960
ctgcttatgt aacaggatgg ggcgctcaag aatatgctgg ccacacagtt ccagagctaa 1020
ggcaaggaca ggtcagaata ataagtaatg atgtatgtaa tgcaccacat agttataatg 1080
gagccatcct gtctggaatg ctgtgtgctg gactaccca aggtggagtg gacgcatgtc 1140
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ggatagtaag ctggggagat cagtgtggcc tgcgggataa gccaggagtg tatactcgag 1260
tgacagcata cattgactgg attaggcaac aaactgggat ctagtgaac aagtgcattc 1320
ctgttgcaaa gtctgtatgc aggtgtgcct gtcttaaat ccaaagcttt acatttcaac 1380
tgaaaaagaa actagaaatg tcctaattta acatcttgtt acataaatat ggtttaacaa 1440
aaaaaaaaa aaaaaactcg ag 1462

<210> 56

<211> 159

<212> PRT

<213> Homo sapiens

<400> 56

Thr Met Tyr Arg Ala Leu Arg Leu Leu Ala Arg Ser Arg Pro Leu Val
1 5 10 15

Arg Ala Pro Ala Ala Ala Leu Ala Ser Ala Pro Gly Leu Gly Gly Ala
20 25 30

Ala Val Pro Ser Phe Trp Pro Pro Asn Ala Ala Arg Met Ala Ser Gln
35 40 45

Asn Ser Phe Arg Ile Glu Tyr Asp Thr Phe Gly Glu Leu Lys Val Pro
50 55 60

Asn Asp Lys Tyr Tyr Gly Ala Gln Thr Val Arg Ser Thr Met Asn Phe
65 70 75 80

Lys Ile Gly Gly Val Thr Glu Arg Met Pro Thr Pro Val Ile Lys Ala
85 90 95

Phe Gly Ile Leu Lys Arg Ala Ala Ala Glu Val Asn Gln Asp Tyr Gly
100 105 110

Leu Asp Pro Lys Ile Ala Asn Ala Ile Met Lys Ala Ala Asp Glu Val
115 120 125

Ala Glu Gly Lys Leu Asn Asp His Phe Pro Leu Val Val Trp Gln Thr
130 135 140

Gly Ser Gly Thr Gln Thr Asn Met Asn Val Asn Glu Val Ile Ser
145 150 155

<210> 57

<211> 165

<212> PRT

<213> Homo sapiens

<400> 57

Lys Lys Ser Met Phe Ala Glu Ile Gln Ile Gln Asp Lys Asp Arg Met
1 5 10 15

Gly Thr Ala Gly Lys Val Ile Lys Cys Lys Ala Ala Val Leu Trp Glu
20 25 30

Gln Lys Gln Pro Phe Ser Ile Glu Glu Ile Glu Val Ala Pro Pro Lys
35 40 45

Thr Lys Glu Val Arg Ile Lys Ile Leu Ala Thr Gly Ile Cys Arg Thr
50 55 60

Asp Asp His Val Ile Lys Gly Thr Met Val Ser Lys Phe Pro Val Ile
65 70 75 80

Val Gly His Glu Ala Thr Gly Ile Val Glu Ser Ile Gly Glu Gly Val
85 90 95

Thr Thr Val Lys Pro Gly Asp Lys Val Ile Pro Leu Phe Leu Pro Gln
100 105 110

Cys Arg Glu Cys Asn Ala Cys Arg Asn Pro Asp Gly Asn Leu Cys Ile
115 120 125

Arg Ser Asp Ile Thr Gly Arg Gly Val Leu Ala Asp Gly Thr Thr Arg
130 135 140

Phe Thr Cys Lys Gly Glu Pro Val His His Phe Met Asn Thr Ser Thr
145 150 155 160

Phe Thr Glu Tyr Thr
165

<210> 58

<211> 259

<212> PRT

<213> Homo sapiens

<400> 58

Glu Ser Glu Gln Lys Gly Lys Ala Ala Leu Ala Ala Thr Leu Glu Glu
 1 5 10 15

Tyr Lys Ala Thr Val Ala Ser Asp Gln Ile Glu Met Asn Arg Leu Lys
 20 25 30

Ala Gln Leu Glu Asn Glu Lys Gln Lys Val Ala Glu Leu Tyr Ser Ile
 35 40 45

His Asn Ser Gly Asp Lys Ser Asp Ile Gln Asp Leu Leu Glu Ser Val
 50 55 60

Arg Leu Asp Lys Glu Lys Ala Glu Thr Leu Ala Ser Ser Leu Gln Glu
 65 70 75 80

Asp Leu Ala His Thr Arg Asn Asp Ala Asn Arg Leu Gln Asp Ala Ile
 85 90 95

Ala Lys Val Glu Asp Glu Tyr Arg Ala Phe Gln Glu Glu Ala Lys Lys
 100 105 110

Gln Ile Glu Asp Leu Asn Met Thr Leu Glu Lys Leu Arg Ser Asp Leu
 115 120 125

Asp Glu Lys Glu Thr Glu Arg Ser Asp Met Lys Glu Thr Ile Phe Glu
 130 135 140

Leu Glu Asp Glu Val Glu Gln His Arg Ala Val Lys Leu His Asp Asn
 145 150 155 160

Leu Ile Ile Ser Asp Leu Glu Asn Thr Val Lys Lys Leu Gln Asp Gln
 165 170 175

Lys His Asp Met Glu Arg Glu Ile Lys Thr Leu His Arg Arg Leu Arg
 180 185 190

Glu Glu Ser Ala Glu Trp Arg Gln Phe Gln Ala Asp Leu Gln Thr Ala
 195 200 205

Val Val Ile Ala Asn Asp Ile Lys Ser Glu Ala Gln Glu Glu Ile Gly
 210 215 220

Asp Leu Lys Arg Arg Leu His Glu Ala Gln Glu Lys Asn Glu Lys Leu
 225 230 235 240

Thr Lys Glu Leu Glu Glu Ile Lys Ser Arg Lys Gln Glu Glu Glu Arg
 245 250 255

Gly Gly Tyr

<210> 59
 <211> 125
 <212> PRT
 <213> Homo sapiens

<400> 59

Gly Thr Ser Phe Ser Lys Asn His Ala Ala Pro Phe Ser Lys Val Leu
 1 5 10 15
 Thr Phe Tyr Arg Lys Glu Pro Phe Thr Leu Glu Ala Tyr Tyr Ser Ser
 20 25 30
 Pro Gln Asp Leu Pro Tyr Pro Asp Pro Ala Ile Ala Gln Phe Ser Val
 35 40 45
 Gln Lys Val Thr Pro Gln Ser Asp Gly Ser Ser Ser Lys Val Lys Val
 50 55 60
 Lys Val Arg Val Asn Val His Gly Ile Phe Ser Val Ser Ser Ala Ser
 65 70 75 80
 Leu Val Glu Val His Lys Ser Glu Glu Asn Glu Glu Pro Met Glu Thr
 85 90 95
 Asp Gln Asn Ala Lys Glu Glu Glu Lys Met Gln Val Asp Gln Glu Glu
 100 105 110
 Pro His Val Glu Glu Gln Gln Gln Gln Thr Pro Gly Arg
 115 120 125

<210> 60
 <211> 246
 <212> PRT
 <213> Homo sapiens

<400> 60

Met Tyr Arg Pro Ala Arg Val Thr Ser Thr Ser Arg Phe Leu Asn Pro
 1 5 10 15
 Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val
 20 25 30
 Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr
 35 40 45
 Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln
 50 55 60
 Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile
 65 70 75 80

Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln
 85 90 95
 Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val
 100 105 110
 Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Asn Gly
 115 120 125
 Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn
 130 135 140
 Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu
 145 150 155 160
 Thr Asp Gln Ala Ala Ala Asn Trp Leu Ile Asn Glu Cys Gly Ala Gly
 165 170 175
 Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu
 180 185 190
 Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn
 195 200 205
 Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr
 210 215 220
 Ala Ala His Cys Phe Arg Ser Asn Ser Asn Pro Arg Asp Trp Ile Ala
 225 230 235 240
 Thr Ser Gly Ile Ser Thr
 245

<210> 61

<211> 128

<212> PRT

<213> Homo sapiens

<400> 61

Gly Ile Phe Ser Val Ser Ser Ala Ser Leu Val Glu Val His Lys Ser
 1 5 10 15

Glu Glu Asn Glu Glu Pro Met Glu Thr Asp Gln Asn Ala Lys Glu Glu
 20 25 30

Glu Lys Met Gln Val Asp Gln Glu Glu Pro His Val Glu Glu Gln Gln
 35 40 45

Gln Gln Thr Pro Ala Glu Asn Lys Ala Glu Ser Glu Glu Met Glu Thr
 50 55 60

Ser Gln Ala Gly Ser Lys Asp Lys Lys Met Asp Gln Pro Pro Gln Ala
 65 70 75 80

Lys Lys Ala Lys Val Lys Thr Ser Thr Val Asp Leu Pro Ile Glu Asn

85 90 95

Gln Leu Leu Trp Gln Ile Asp Arg Glu Met Leu Asn Leu Tyr Ile Glu
100 105 110

Asn Glu Gly Lys Met Ile Met Gln Asp Lys Leu Glu Lys Glu Arg Asn
115 120 125

<210> 62
 <211> 418
 <212> PRT
 <213> Homo sapiens

<400> 62
 Met Tyr Arg Pro Ala Arg Val Thr Ser Thr Ser Arg Phe Leu Asn Pro
1 5 10 15

Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val
20 25 30

Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr
35 40 45

Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln
50 55 60

Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile
65 70 75 80

Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln
85 90 95

Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val
100 105 110

Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Asn Gly
115 120 125

Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn
130 135 140

Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu
145 150 155 160

Thr Asp Gln Ala Ala Ala Asn Trp Leu Ile Asn Glu Cys Gly Ala Gly
165 170 175

Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu
180 185 190

Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn
195 200 205

Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr

210 215 220
 Ala Ala His Cys Phe Arg Ser Asn Ser Asn Pro Arg Asp Trp Ile Ala
 225 230 235 240
 Thr Ser Gly Ile Ser Thr Thr Phe Pro Lys Leu Arg Met Arg Val Arg
 245 250 255
 Asn Ile Leu Ile His Asn Asn Tyr Lys Ser Ala Thr His Glu Asn Asp
 260 265 270
 Ile Ala Leu Val Arg Leu Glu Asn Ser Val Thr Phe Thr Lys Asp Ile
 275 280 285
 His Ser Val Cys Leu Pro Ala Ala Thr Gln Asn Ile Pro Pro Gly Ser
 290 295 300
 Thr Ala Tyr Val Thr Gly Trp Gly Ala Gln Glu Tyr Ala Gly His Thr
 305 310 315 320
 Val Pro Glu Leu Arg Gln Gly Gln Val Arg Ile Ile Ser Asn Asp Val
 325 330 335
 Cys Asn Ala Pro His Ser Tyr Asn Gly Ala Ile Leu Ser Gly Met Leu
 340 345 350
 Cys Ala Gly Val Pro Gln Gly Gly Val Asp Ala Cys Gln Gly Asp Ser
 355 360 365
 Gly Gly Pro Leu Val Gln Glu Asp Ser Arg Arg Leu Trp Phe Ile Val
 370 375 380
 Gly Ile Val Ser Trp Gly Asp Gln Cys Gly Leu Pro Asp Lys Pro Gly
 385 390 395 400
 Val Tyr Thr Arg Val Thr Ala Tyr Ile Asp Trp Ile Arg Gln Gln Thr
 405 410 415

Gly Ile

<210> 63

<211> 776

<212> DNA

<213> Homo sapiens

<400> 63

cacagatggt gatagaggaa tccatcttgc agtcagataa agccctcact gatagagaga 60
 aggcagtagc agtggatcgg gccaaagaagg aggcagctga gaaggaacag gaacttttaa 120
 aacagaaatt acaggagcag ccagcaacag atggaggctc aagataagag tcgcaaggaa 180
 aactagccaa ctgaaggaga agctgcagat ggagagagaa cacctactga gagagcagat 240
 tatgatgttg gagcacacgc agaagggtcca aaatgattgg cttcatgaag gatttaagaa 300
 gaagtatgag gagatgaatg cagagataag tcaatttaaa cgtatgattg atactacaaa 360
 aaatgatgat actccctgga ttgcacgaac cttggacaac cttgccgatg agctaactgc 420
 aatattgtct gctcctgcta aattaattgg tcatgggtgtc aaagggtgtga gctcactctt 480

taaaaagcat aagctcccct tttaaggata ttatagattg tacatatatg ctttggacta 540
 tttttgatct gtatgttttt cattttcatt cagcaagttt tttttttttt tcagagtctt 600
 actctgttgc ccaggctgga gtacagtggg gcaatctcag ctactgcaa cctctgcctc 660
 ctgggttcaa gagattcacc tgcctcagcc ccctagtagc tgggattata ggtgtacacc 720
 accacaccca gctaattttt gtatttttag tagagatggg gtttcactat gttggc 776

<210> 64

<211> 160

<212> DNA

<213> Homo sapiens

<400> 64

gcagcgctct cggttgcagt acccactgga aggacttagg cgctcgctg gacaccgcaa 60
 gccctcagt agcctcggcc caagaggcct gctttccact cgctagcccc gccgggggtc 120
 cgtgtcctgt ctcggtggcc ggaccggggc ccgagcccca 160

<210> 65

<211> 72

<212> PRT

<213> Homo sapiens

<400> 65

Leu Ser Ala Met Gly Phe Thr Ala Ala Gly Ile Ala Ser Ser Ser Ile
 1 5 10 15

Ala Ala Lys Met Met Ser Ala Ala Ala Ile Ala Asn Gly Gly Gly Val
 20 25 30

Ala Ser Gly Ser Leu Val Ala Thr Leu Gln Ser Leu Gly Ala Thr Gly
 35 40 45

Leu Ser Gly Leu Thr Lys Phe Ile Leu Gly Ser Ile Gly Ser Ala Ile
 50 55 60

Ala Ala Val Ile Ala Arg Phe Tyr
 65 70

<210> 66

<211> 2581

<212> DNA

<213> Homo sapiens

<400> 66

ctttcaaccc gcgctcgccg gctccagccc cgcgcgcccc cacccttgc cctcccgggc 60
 gctccgcagg gtgaggtggc tttgacccc gggtgcccgc ccagcacgac cgaggaggtg 120
 gctggaacgc tggaggatga acggagaagc cgactgcccc acagacctgg aaatggccgc 180
 ccccaaaggc caagaccgtt ggtcccagga agacatgctg actttgctgg aatgcatgaa 240
 gaacaacctt ccatccaatg acagctccaa gttcaaaacc accgaatcac acatggactg 300
 ggaaaaagta gcatttaaag acttttctgg agacatgtgc aagctcaa at ggtggagat 360
 ttctaataag gtgaggaagt tccgtacatt gacagaattg atcctcgatg ctgaggaaca 420
 tgttaaaaat ctttacaag gcaaaaaact caagaaacac ccagacttcc caaagaagcc 480
 cctgacccct tatttccgct tcttcatgga gaagcggggc aagtatgca aactccaccc 540
 tgagatgagc aacctggacc taaccaagat tctgtccaag aaatacaagg agcttccgga 600
 gaagaagaag atgaaatata ttcaggactt ccagagagag aaacaggagt tcgagcgaaa 660

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cctggccccga ttcaggggagg atcacccccga cctaattccag aatgcccaaga aatcggacat 720
cccagagaag cccaaaaccc cccagcagct gtggtacacc cagcagaaga aggtgtatct 780
caaagtgcgg ccagatgccca ctacgaagga ggtgaaggac tccctgggga agcagtggtc 840
tcagctctcg gacaaaaaga ggctgaaatg gattcataag gccctggagc agcgggaagga 900
gtacgaggag atcatgagag actatatcca gaagcaccca gagctgaaca tcagtgagga 960
gggtatcacc aagtccaccc tcaccaaggc cgaacgccag ctcaaggaca agtttgacgg 1020
gcgacccacc aagccacctc cgaacagcta ctgcgtgtac tgcgcagagc tcattggccaa 1080
catgaaggag gtgcccagca cagagcgcac ggtgctgtgc agccagcagt ggaagctgct 1140
gtcccagaag gagaaggacg cctatcacaa gaagtgtgat cagaaaaaga aagattacga 1200
ggtggagctg ctccgtttcc tcgagagcct gcctgaggag gagcagcagc gggctctggg 1260
ggaagagaag atgctgaaca tcaacaagaa gcaggccacc agccccgcct ccaagaagcc 1320
agcccaggaa gggggcaagg gcggctccga gaagcccaag cgggccgtgt cggccatgtt 1380
catcttctcg gaggagaaac ggcggcagct gcaggaggag cggcctgagc tctccgagag 1440
cgagctgacc cgcctgctgg cccgaatgtg gaacgacctg tctgagaaga agaaggccaa 1500
gtacaaggcc cgagaggcgg cgctcaaggc tcagtcggag aggaagcccg gcggggagcg 1560
cgaggaacgg ggcaagctgc ccgagtcacc caaaagagct gaggagatct ggcaacagag 1620
cgttatcggc gactacctgg ccgcgttcaa gaatgaccgg gtgaaggcct tgaaagccat 1680
ggaaatgacc tggaataaca tggaaaagaa ggagaaactg atgtggatta agaaggcagc 1740
cgaagaccaa aagcgatatg agagagagct gagtgaagat cgggcacctc cagctgctac 1800
aaattcttcc aagaagatga aattccaggg agaacccaag aagcctccca tgaacggtta 1860
ccagaagttc tcccaggagc tgctgtccaa tggggagctg aaccacctgc cgctgaagga 1920
gcgcatggtg gagatcgcca gtgcgtggca gcgcattctc cagagccaga aggagcacta 1980
caaaaagctg gccgaggagc agcaaaagca gtacaagggtg cacctggacc tctgggttaa 2040
gagcctgtct ccccaggacc gtgcagcata taaagagtac atctccaata aacgtaagag 2100
catgaccaag ctgcgaggcc caaaccacca atccagccgg actactctgc agtccaagtc 2160
ggagtccgag gaggatgatg aagaggatga ggatgacgag gacgaggatg aagaagagga 2220
agatgatgag aatggggact cctctgaaga tggcggcgac tcctctgagt ccagcagcga 2280
ggacgagagc gaggatgggg atgagaatga agaggatgac gaggacgaag acgacgacga 2340
ggatgacgat gaggatgaag ataatgagtc cgagggcagc agctccagct cctcctcctt 2400
aggggactcc tcagactttg actccaactg aggcttagcc ccacccagg ggagccaggg 2460
agagcccagg agtccccctc cccaactgac cacctttgtt tcttccccat gttctgtccc 2520
ttgccccctt ggccctcccc actttctttc tttctttaa aaaaaaaaaa aaaaactcga 2580
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<210> 67

<211> 764

<212> PRT

<213> Homo sapiens

<400> 67

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Met Asn Gly Glu Ala Asp Cys Pro Thr Asp Leu Glu Met Ala Ala Pro
  1                      5                      10                      15

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Lys Gly Gln Asp Arg Trp Ser Gln Glu Asp Met Leu Thr Leu Leu Glu
      20                      25                      30

```

```

Cys Met Lys Asn Asn Leu Pro Ser Asn Asp Ser Ser Lys Phe Lys Thr
      35                      40                      45

```

```

Thr Glu Ser His Met Asp Trp Glu Lys Val Ala Phe Lys Asp Phe Ser
      50                      55                      60

```

```

Gly Asp Met Cys Lys Leu Lys Trp Val Glu Ile Ser Asn Glu Val Arg
      65                      70                      75                      80

```

Lys Phe Arg Thr Leu Thr Glu Leu Ile Leu Asp Ala Gln Glu His Val
 85 90 95
 Lys Asn Pro Tyr Lys Gly Lys Lys Leu Lys Lys His Pro Asp Phe Pro
 100 105 110
 Lys Lys Pro Leu Thr Pro Tyr Phe Arg Phe Phe Met Glu Lys Arg Ala
 115 120 125
 Lys Tyr Ala Lys Leu His Pro Glu Met Ser Asn Leu Asp Leu Thr Lys
 130 135 140
 Ile Leu Ser Lys Lys Tyr Lys Glu Leu Pro Glu Lys Lys Lys Met Lys
 145 150 155 160
 Tyr Ile Gln Asp Phe Gln Arg Glu Lys Gln Glu Phe Glu Arg Asn Leu
 165 170 175
 Ala Arg Phe Arg Glu Asp His Pro Asp Leu Ile Gln Asn Ala Lys Lys
 180 185 190
 Ser Asp Ile Pro Glu Lys Pro Lys Thr Pro Gln Gln Leu Trp Tyr Thr
 195 200 205
 His Glu Lys Lys Val Tyr Leu Lys Val Arg Pro Asp Ala Thr Thr Lys
 210 215 220
 Glu Val Lys Asp Ser Leu Gly Lys Gln Trp Ser Gln Leu Ser Asp Lys
 225 230 235 240
 Lys Arg Leu Lys Trp Ile His Lys Ala Leu Glu Gln Arg Lys Glu Tyr
 245 250 255
 Glu Glu Ile Met Arg Asp Tyr Ile Gln Lys His Pro Glu Leu Asn Ile
 260 265 270
 Ser Glu Glu Gly Ile Thr Lys Ser Thr Leu Thr Lys Ala Glu Arg Gln
 275 280 285
 Leu Lys Asp Lys Phe Asp Gly Arg Pro Thr Lys Pro Pro Pro Asn Ser
 290 295 300
 Tyr Ser Leu Tyr Cys Ala Glu Leu Met Ala Asn Met Lys Asp Val Pro
 305 310 315 320
 Ser Thr Glu Arg Met Val Leu Cys Ser Gln Gln Trp Lys Leu Leu Ser
 325 330 335
 Gln Lys Glu Lys Asp Ala Tyr His Lys Lys Cys Asp Gln Lys Lys Lys
 340 345 350
 Asp Tyr Glu Val Glu Leu Leu Arg Phe Leu Glu Ser Leu Pro Glu Glu
 355 360 365
 Glu Gln Gln Arg Val Leu Gly Glu Glu Lys Met Leu Asn Ile Asn Lys

370 375 380
 Lys Gln Ala Thr Ser Pro Ala Ser Lys Lys Pro Ala Gln Glu Gly Gly
 385 390 395 400
 Lys Gly Gly Ser Glu Lys Pro Lys Arg Pro Val Ser Ala Met Phe Ile
 405 410 415
 Phe Ser Glu Glu Lys Arg Arg Gln Leu Gln Glu Glu Arg Pro Glu Leu
 420 425 430
 Ser Glu Ser Glu Leu Thr Arg Leu Leu Ala Arg Met Trp Asn Asp Leu
 435 440 445
 Ser Glu Lys Lys Lys Ala Lys Tyr Lys Ala Arg Glu Ala Ala Leu Lys
 450 455 460
 Ala Gln Ser Glu Arg Lys Pro Gly Gly Glu Arg Glu Glu Arg Gly Lys
 465 470 475 480
 Leu Pro Glu Ser Pro Lys Arg Ala Glu Glu Ile Trp Gln Gln Ser Val
 485 490 495
 Ile Gly Asp Tyr Leu Ala Arg Phe Lys Asn Asp Arg Val Lys Ala Leu
 500 505 510
 Lys Ala Met Glu Met Thr Trp Asn Asn Met Glu Lys Lys Glu Lys Leu
 515 520 525
 Met Trp Ile Lys Lys Ala Ala Glu Asp Gln Lys Arg Tyr Glu Arg Glu
 530 535 540
 Leu Ser Glu Met Arg Ala Pro Pro Ala Ala Thr Asn Ser Ser Lys Lys
 545 550 555 560
 Met Lys Phe Gln Gly Glu Pro Lys Lys Pro Pro Met Asn Gly Tyr Gln
 565 570 575
 Lys Phe Ser Gln Glu Leu Leu Ser Asn Gly Glu Leu Asn His Leu Pro
 580 585 590
 Leu Lys Glu Arg Met Val Glu Ile Gly Ser Arg Trp Gln Arg Ile Ser
 595 600 605
 Gln Ser Gln Lys Glu His Tyr Lys Lys Leu Ala Glu Glu Gln Gln Lys
 610 615 620
 Gln Tyr Lys Val His Leu Asp Leu Trp Val Lys Ser Leu Ser Pro Gln
 625 630 635 640
 Asp Arg Ala Ala Tyr Lys Glu Tyr Ile Ser Asn Lys Arg Lys Ser Met
 645 650 655
 Thr Lys Leu Arg Gly Pro Asn Pro Lys Ser Ser Arg Thr Thr Leu Gln
 660 665 670

Ser Lys Ser Glu Ser Glu Glu Asp Asp Glu Glu Asp Glu Asp Asp Glu
 675 680 685
 Asp Glu Asp Glu Glu Glu Glu Asp Asp Glu Asn Gly Asp Ser Ser Glu
 690 695 700
 Asp Gly Gly Asp Ser Ser Glu Ser Ser Ser Glu Asp Glu Ser Glu Asp
 705 710 715 720
 Gly Asp Glu Asn Glu Glu Asp Asp Glu Asp Glu Asp Asp Asp Glu Asp
 725 730 735
 Asp Asp Glu Asp Glu Asp Asn Glu Ser Glu Gly Ser Ser Ser Ser Ser
 740 745 750
 Ser Ser Leu Gly Asp Ser Ser Asp Phe Asp Ser Asn
 755 760

<210> 68
 <211> 434
 <212> DNA
 <213> Homo sapiens

<400> 68
 ctaagatgct ggatgctgaa gacatcgctc gaactgcccg gccagatgag aaagccatta 60
 tgacttatgt gtctagcttc tatcatgcct tctctggagc ccagaaggca gaaacagcag 120
 ccaatcgcat ctgcaaagtg ttggcgggtca atcaagagaa cgagcagctt atggaagact 180
 atgagaagct ggccagtgat ctgttggagt ggatccgccg caccatccca tggctggaga 240
 atcgggtgcc tgagaacacc atgcatgccca tgcagcagaa gctggaggac ttccgagact 300
 atagacgcct gcacaagccg cccaagggtgc aggagaagtg ccagctggag atcaacttta 360
 acacgctgca gaccaaactg cggctcagca accggcctgc cttcatgccc tccgagggca 420
 ggatgggtctc ggat 434

<210> 69
 <211> 244
 <212> DNA
 <213> Homo sapiens

<400> 69
 aggcagcatg ctcggtgaga gtcatacaca ctccctaata tcaagtacgc agggacacaa 60
 aactgcgga aggccgcagg gtcctctgcc taggaaaacc agagaccttt gtccacttgt 120
 ttatgtgctg accttccctc cactattgtc ctgtgaccct gccaaatccc cttttgtgag 180
 aaacacccaa gaatgatcaa taaaaaataa attaatttag gaaaaaaaaa aaaaaaaact 240
 cgag 244

<210> 70
 <211> 437
 <212> DNA
 <213> Homo sapiens

<400> 70
 ctgggacggg agcgtccagc gggactcgaa cccagatgt gaaggcgttt ctggaaagtc 60
 cttggtccct ggatccagcg tcggccagcc cagagcccgt gccgcacatc cttgcgtcct 120

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ccaggcagtg ggacccccg agctgcacgt ccctgggcac ggacaagtgt gaggcactgt 180
tggggctgtg ccagggtgcgg ggtgggctgc cccctttctc agaacttcc agcctgggtgc 240
cgtggcccc aggccggagt ctctctaagg ctgtgaggcc acccctgtcc tggcctccgt 300
tctcgcagca gcagacctg cccgtgatga gcggggaggc ccttggctgg ctggggccagg 360
ctggttcctt ggccatgggg gctgcacctc tgggggagcc agccaaggag gaccccatgc 420
tggcgcagga agccggg

```

<210> 71

<211> 271

<212> DNA

<213> Homo sapiens

<400> 71

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gcgcagagtt ctgtcgtcca ccatcgagt aggaagagag cattgggtcc cctgagatag 60
aagagatggc tctcttcagt gccagtcctc catacattaa cccgatcatc ccccttactg 120
gaccaatcca aggagggctg caggagggac ttcaggtgac cctccagggg actaccgaga 180
gttttgcaca aaagtttgtg gtgaactttt cagaacagct tcaatggaga tgacttgcc 240
ttccacttca accccgggta tgaggagga g

```

<210> 72

<211> 290

<212> DNA

<213> Homo sapiens

<400> 72

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ccgagcccta cccggaggtc tccagaatcc ccaccgtcag gggatgcaac ggctccctgt 60
ctggtgccct ctctgctgc gaggactcgg cccagggctc gggcccgccc aaggccccta 120
cgggtggccga ggggtcccagc tctgccttc ggcggaacgt gatcagcgag agggagcgca 180
ggaagcggat gtcgttgagc tgtgagcgtc tgcgggccct gctgccccag ttcgatggcc 240
ggcgggagga catggcctcg gtcctggaga tgtctgttgc aattcctgcg

```

<210> 73

<211> 144

<212> PRT

<213> Homo sapiens

<400> 73

```

Lys Met Leu Asp Ala Glu Asp Ile Val Gly Thr Ala Arg Pro Asp Glu
  1             5             10             15

```

```

Lys Ala Ile Met Thr Tyr Val Ser Ser Phe Tyr His Ala Phe Ser Gly
          20             25             30

```

```

Ala Gln Lys Ala Glu Thr Ala Ala Asn Arg Ile Cys Lys Val Leu Ala
          35             40             45

```

```

Val Asn Gln Glu Asn Glu Gln Leu Met Glu Asp Tyr Glu Lys Leu Ala
          50             55             60

```

```

Ser Asp Leu Leu Glu Trp Ile Arg Arg Thr Ile Pro Trp Leu Glu Asn
          65             70             75             80

```

```

Arg Val Pro Glu Asn Thr Met His Ala Met Gln Gln Lys Leu Glu Asp
          85             90             95

```

Phe Arg Asp Tyr Arg Arg Leu His Lys Pro Pro Lys Val Gln Glu Lys
 100 105 110

Cys Gln Leu Glu Ile Asn Phe Asn Thr Leu Gln Thr Lys Leu Arg Leu
 115 120 125

Ser Asn Arg Pro Ala Phe Met Pro Ser Glu Gly Arg Met Val Ser Asp
 130 135 140

<210> 74

<211> 64

<212> PRT

<213> Homo sapiens

<400> 74

Gly Ser Met Leu Val Glu Ser His His His Ser Leu Ile Ser Ser Thr
 1 5 10 15

Gln Gly His Lys His Cys Gly Arg Pro Gln Gly Pro Leu Pro Arg Lys
 20 25 30

Thr Arg Asp Leu Cys Ser Leu Val Tyr Val Leu Thr Phe Pro Pro Leu
 35 40 45

Leu Ser Cys Asp Pro Ala Lys Ser Pro Phe Val Arg Asn Thr Gln Glu
 50 55 60

<210> 75

<211> 145

<212> PRT

<213> Homo sapiens

<400> 75

Gly Thr Gly Ala Ser Ser Gly Thr Arg Thr Pro Asp Val Lys Ala Phe
 1 5 10 15

Leu Glu Ser Pro Trp Ser Leu Asp Pro Ala Ser Ala Ser Pro Glu Pro
 20 25 30

Val Pro His Ile Leu Ala Ser Ser Arg Gln Trp Asp Pro Ala Ser Cys
 35 40 45

Thr Ser Leu Gly Thr Asp Lys Cys Glu Ala Leu Leu Gly Leu Cys Gln
 50 55 60

Val Arg Gly Gly Leu Pro Pro Phe Ser Glu Pro Ser Ser Leu Val Pro
 65 70 75 80

Trp Pro Pro Gly Arg Ser Leu Pro Lys Ala Val Arg Pro Pro Leu Ser
 85 90 95

Trp Pro Pro Phe Ser Gln Gln Gln Thr Leu Pro Val Met Ser Gly Glu
 100 105 110

Ala Leu Gly Trp Leu Gly Gln Ala Gly Ser Leu Ala Met Gly Ala Ala
 115 120 125

Pro Leu Gly Glu Pro Ala Lys Glu Asp Pro Met Leu Ala Gln Glu Ala
 130 135 140

Gly
 145

<210> 76

<211> 69

<212> PRT

<213> Homo sapiens

<400> 76

Ala Glu Phe Cys Arg Pro Pro Ser Ser Glu Glu Glu Ser Ile Gly Ser
 1 5 10 15

Pro Glu Ile Glu Glu Met Ala Leu Phe Ser Ala Gln Ser Pro Tyr Ile
 20 25 30

Asn Pro Ile Ile Pro Phe Thr Gly Pro Ile Gln Gly Gly Leu Gln Glu
 35 40 45

Gly Leu Gln Val Thr Leu Gln Gly Thr Thr Glu Ser Phe Ala Gln Lys
 50 55 60

Phe Val Val Asn Phe
 65

<210> 77

<211> 96

<212> PRT

<213> Homo sapiens

<400> 77

Glu Pro Tyr Pro Glu Val Ser Arg Ile Pro Thr Val Arg Gly Cys Asn
 1 5 10 15

Gly Ser Leu Ser Gly Ala Leu Ser Cys Cys Glu Asp Ser Ala Gln Gly
 20 25 30

Ser Gly Pro Pro Lys Ala Pro Thr Val Ala Glu Gly Pro Ser Ser Cys
 35 40 45

Leu Arg Arg Asn Val Ile Ser Glu Arg Glu Arg Arg Lys Arg Met Ser
 50 55 60

Leu Ser Cys Glu Arg Leu Arg Ala Leu Leu Pro Gln Phe Asp Gly Arg
 65 70 75 80

Arg Glu Asp Met Ala Ser Val Leu Glu Met Ser Val Ala Ile Pro Ala

85

90

95

<210> 78
 <211> 2076
 <212> DNA
 <213> Homo sapiens

<400> 78
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 gaaaagttat taaatgcaaa gcagctgtgc tttgggagca gaagcaaccc ttctccattg 120
 aggaaataga agttgcccc ccaaagacta aagaagtctg cattaagatt ttggccacag 180
 gaatctgtcg cacagatgac catgtgataa aaggaacaat ggtgtccaag tttccagtga 240
 ttgtgggaca tgaggcaact gggattgtag agagcattgg agaaggagt actacagtga 300
 aaccaggtga caaagtcac cctctcttct tgcacacaatg tagagaatgc aatgcttgct 360
 gcaaccaga tggcaacctt tgcattagga gcgatattac tggctgtgga gtactggctg 420
 atggcaccac cagatttaca tgcaagggca aaccagtcca ccacttcacg aacaccagta 480
 catttaccga gtacacagtg gtggatgaat cttctgttgc taagattgat gatgcagctc 540
 ctctgagaa agtctgttta attggctgtg ggttttccac tggatatggc gctgctgtta 600
 aaactggcaa ggtcaaacct ggttccactt gcgtcgtctt tggcctgaga ggagtggcc 660
 tgtcagtcac catgggctgt aagtcagctg gtgcattctag gatcattggg attgacctca 720
 acaaagacaa atttgagaag gccatggctg taggtgccac tgagtgtatc agtcccaagg 780
 actctacca acccatcagt gaggtgctgt cagaaatgac aggcaacaac gtgggatata 840
 cctttgaagt tattgggcat cttgaaacca tgattgatgc cctggcatcc tgccacatga 900
 actatgggac cagcgtgggt gtaggagttc ctccatcagc caagatgctc acctatgacc 960
 cgatgttgct cttcactgga cgcacatgga agggatgtgt ctttggagggt ttgaaaagca 1020
 gagatgatgt cccaaaacta gtgactgagt tcctggcaaa gaaatttgac ctggaccagt 1080
 tgataactca tgtcttacca tttaaaaaaa tcagtgaagg atttgagctg ctcaattcag 1140
 gacaaagcat tcgaacgggt ctgacgtttt gagatccaaa gtggcaggag gtctgtgttg 1200
 tcatggtgaa ctggagtttc tctgtgaga gttccctcat ctgaaatcat gtatctgtct 1260
 cacaataca agcataagta gaagatttgc tgaagacata gaaccttat aaagaattat 1320
 taacctttat aaacatttaa agtcttgcga gcacctggga attagtataa taacaatgtt 1380
 aatatttttg atttacattt tgtaaggcta taattgtatc ttttaagaaa acatacattt 1440
 ggattttctat gttgaaatgg agatttttaa gagttttaac cagctgctgc agatatatat 1500
 ctcaaaacag atatagcgta taaagatata gtaaagcat ctctagagt aatattcact 1560
 taacacattg aaactattat tttttagatt tgaatataaa tgtatttttt aaacacttgt 1620
 tatgagttaa cttggattac attttgaaat cagttcattc catgatgcat attactggat 1680
 tagattaaga aagacagaaa agattaaggg acgggcacat ttttcaacga ttaagaatca 1740
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 ttaagtcctt attcactgtg cttagtagtg actccattta ataaaaagtg ttttagttt 1980
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<210> 79
 <211> 2790
 <212> DNA
 <213> Homo sapiens

<400> 79
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 caggggtagt gatcctggca gtcaccatag ctctacttgt ttacttttta gcttttgcac 180
 aaaaatctta cttttatagg agcagttttc aactcctaaa tgttgaatat aatagtcagt 240

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taaattcacc agctacacag gaatacagga ctttgagtgg aagaattgaa tctctgatta 300
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tgaggcaaga tggtagtggg gtgagagcgg atgttgtcat gaaatttcaa ttcactagaa 420
ataacaatgg agcatcaatg aaaagcagaa ttgagtctgt tttagcacia atgctgaata 480
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```

<210> 80

<211> 1460

<212> DNA

<213> Homo sapiens

<400> 80

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gtcgcagggg tagtgatcct ggcagtcacc atagctctac ttgtttactt tttagctttt 180
gatcaaaaat cttactttta taggagcagt tttcaactcc taaatgttga atataatag 240
cagttaaatt caccagctac acaggaatac aggactttga gtggaagaat tgaatctctg 300

```

```

attactaaaa cattcaaaga atcaaattta agaaatcagt tcatcagagc tcatgttgcc 360
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agaaataaca atggagcatc aatgaaaagc agaattgagt ctgttttacg acaaagtctg 480
aataactctg gaaacctgga aataaaccct tcaactgaga taacatcact tactgaccag 540
gctgcagcaa attggcttat taatgaatgt ggggccggtc cagacctaat aacattgtct 600
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acaaaaaaaa aaaaaaaaaa 1460

```

<210> 81

<211> 386

<212> PRT

<213> Homo sapiens

<400> 81

```

Met Phe Ala Glu Ile Gln Ile Gln Asp Lys Asp Arg Met Gly Thr Ala
  1             5             10             15

```

```

Gly Lys Val Ile Lys Cys Lys Ala Ala Val Leu Trp Glu Gln Lys Gln
      20             25             30

```

```

Pro Phe Ser Ile Glu Glu Ile Glu Val Ala Pro Pro Lys Thr Lys Glu
      35             40             45

```

```

Val Arg Ile Lys Ile Leu Ala Thr Gly Ile Cys Arg Thr Asp Asp His
      50             55             60

```

```

Val Ile Lys Gly Thr Met Val Ser Lys Phe Pro Val Ile Val Gly His
      65             70             75             80

```

```

Glu Ala Thr Gly Ile Val Glu Ser Ile Gly Glu Gly Val Thr Thr Val
      85             90             95

```

```

Lys Pro Gly Asp Lys Val Ile Pro Leu Phe Leu Pro Gln Cys Arg Glu
     100             105             110

```

```

Cys Asn Ala Cys Arg Asn Pro Asp Gly Asn Leu Cys Ile Arg Ser Asp
     115             120             125

```

```

Ile Thr Gly Arg Gly Val Leu Ala Asp Gly Thr Thr Arg Phe Thr Cys
     130             135             140

```

```

Lys Gly Lys Pro Val His His Phe Met Asn Thr Ser Thr Phe Thr Glu

```


145 150 155 160
 Tyr Thr Val Val Asp Glu Ser Ser Val Ala Lys Ile Asp Asp Ala Ala
 165 170 175
 Pro Pro Glu Lys Val Cys Leu Ile Gly Cys Gly Phe Ser Thr Gly Tyr
 180 185 190
 Gly Ala Ala Val Lys Thr Gly Lys Val Lys Pro Gly Ser Thr Cys Val
 195 200 205
 Val Phe Gly Leu Arg Gly Val Gly Leu Ser Val Ile Met Gly Cys Lys
 210 215 220
 Ser Ala Gly Ala Ser Arg Ile Ile Gly Ile Asp Leu Asn Lys Asp Lys
 225 230 235 240
 Phe Glu Lys Ala Met Ala Val Gly Ala Thr Glu Cys Ile Ser Pro Lys
 245 250 255
 Asp Ser Thr Lys Pro Ile Ser Glu Val Leu Ser Glu Met Thr Gly Asn
 260 265 270
 Asn Val Gly Tyr Thr Phe Glu Val Ile Gly His Leu Glu Thr Met Ile
 275 280 285
 Asp Ala Leu Ala Ser Cys His Met Asn Tyr Gly Thr Ser Val Val Val
 290 295 300
 Gly Val Pro Pro Ser Ala Lys Met Leu Thr Tyr Asp Pro Met Leu Leu
 305 310 315 320
 Phe Thr Gly Arg Thr Trp Lys Gly Cys Val Phe Gly Gly Leu Lys Ser
 325 330 335
 Arg Asp Asp Val Pro Lys Leu Val Thr Glu Phe Leu Ala Lys Lys Phe
 340 345 350
 Asp Leu Asp Gln Leu Ile Thr His Val Leu Pro Phe Lys Lys Ile Ser
 355 360 365
 Glu Gly Phe Glu Leu Leu Asn Ser Gly Gln Ser Ile Arg Thr Val Leu
 370 375 380
 Thr Phe
 385

<210> 82

<211> 418

<212> PRT

<213> Homo sapiens

<400> 82

Met Tyr Arg Pro Ala Arg Val Thr Ser Thr Ser Arg Phe Leu Asn Pro

1	5	10	15
Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val	20	25	30
Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr	35	40	45
Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln	50	55	60
Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile	65	70	75
Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln	85	90	95
Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val	100	105	110
Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Asn Gly	115	120	125
Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn	130	135	140
Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu	145	150	155
Thr Asp Gln Ala Ala Ala Asn Trp Leu Ile Asn Glu Cys Gly Ala Gly	165	170	175
Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu	180	185	190
Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn	195	200	205
Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr	210	215	220
Ala Ala His Cys Phe Arg Ser Asn Ser Asn Pro Arg Asp Trp Ile Ala	225	230	235
Thr Ser Gly Ile Ser Thr Thr Phe Pro Lys Leu Arg Met Arg Val Arg	245	250	255
Asn Ile Leu Ile His Asn Asn Tyr Lys Ser Ala Thr His Glu Asn Asp	260	265	270
Ile Ala Leu Val Arg Leu Glu Asn Ser Val Thr Phe Thr Lys Asp Ile	275	280	285
His Ser Val Cys Leu Pro Ala Ala Thr Gln Asn Ile Pro Pro Gly Ser	290	295	300

Thr Ala Tyr Val Thr Gly Trp Gly Ala Gln Glu Tyr Ala Gly His Thr
 305 310 315 320
 Val Pro Glu Leu Arg Gln Gly Gln Val Arg Ile Ile Ser Asn Asp Val
 325 330 335
 Cys Asn Ala Pro His Ser Tyr Asn Gly Ala Ile Leu Ser Gly Met Leu
 340 345 350
 Cys Ala Gly Val Pro Gln Gly Gly Val Asp Ala Cys Gln Gly Asp Ser
 355 360 365
 Gly Gly Pro Leu Val Gln Glu Asp Ser Arg Arg Leu Trp Phe Ile Val
 370 375 380
 Gly Ile Val Ser Trp Gly Asp Gln Cys Gly Leu Pro Asp Lys Pro Gly
 385 390 395 400
 Val Tyr Thr Arg Val Thr Ala Tyr Leu Asp Trp Ile Arg Gln Gln Thr
 405 410 415
 Gly Ile

<210> 83
 <211> 418
 <212> PRT
 <213> Homo sapiens

<400> 83

Met Tyr Arg Pro Ala Arg Val Thr Ser Thr Ser Arg Phe Leu Asn Pro
 1 5 10 15
 Tyr Val Val Cys Phe Ile Val Val Ala Gly Val Val Ile Leu Ala Val
 20 25 30
 Thr Ile Ala Leu Leu Val Tyr Phe Leu Ala Phe Asp Gln Lys Ser Tyr
 35 40 45
 Phe Tyr Arg Ser Ser Phe Gln Leu Leu Asn Val Glu Tyr Asn Ser Gln
 50 55 60
 Leu Asn Ser Pro Ala Thr Gln Glu Tyr Arg Thr Leu Ser Gly Arg Ile
 65 70 75 80
 Glu Ser Leu Ile Thr Lys Thr Phe Lys Glu Ser Asn Leu Arg Asn Gln
 85 90 95
 Phe Ile Arg Ala His Val Ala Lys Leu Arg Gln Asp Gly Ser Gly Val
 100 105 110
 Arg Ala Asp Val Val Met Lys Phe Gln Phe Thr Arg Asn Asn Asn Gly
 115 120 125

Ala Ser Met Lys Ser Arg Ile Glu Ser Val Leu Arg Gln Met Leu Asn
 130 135 140
 Asn Ser Gly Asn Leu Glu Ile Asn Pro Ser Thr Glu Ile Thr Ser Leu
 145 150 155 160
 Thr Asp Gln Ala Ala Ala Asn Trp Leu Ile Asn Glu Cys Gly Ala Gly
 165 170 175
 Pro Asp Leu Ile Thr Leu Ser Glu Gln Arg Ile Leu Gly Gly Thr Glu
 180 185 190
 Ala Glu Glu Gly Ser Trp Pro Trp Gln Val Ser Leu Arg Leu Asn Asn
 195 200 205
 Ala His His Cys Gly Gly Ser Leu Ile Asn Asn Met Trp Ile Leu Thr
 210 215 220
 Ala Ala His Cys Phe Arg Ser Asn Ser Asn Pro Arg Asp Trp Ile Ala
 225 230 235 240
 Thr Ser Gly Ile Ser Thr Thr Phe Pro Lys Leu Arg Met Arg Val Arg
 245 250 255
 Asn Ile Leu Ile His Asn Asn Tyr Lys Ser Ala Thr His Glu Asn Asp
 260 265 270
 Ile Ala Leu Val Arg Leu Glu Asn Ser Val Thr Phe Thr Lys Asp Ile
 275 280 285
 His Ser Val Cys Leu Pro Ala Ala Thr Gln Asn Ile Pro Pro Gly Ser
 290 295 300
 Thr Ala Tyr Val Thr Gly Trp Gly Ala Gln Glu Tyr Ala Gly His Thr
 305 310 315 320
 Val Pro Glu Leu Arg Gln Gly Gln Val Arg Ile Ile Ser Asn Asp Val
 325 330 335
 Cys Asn Ala Pro His Ser Tyr Asn Gly Ala Ile Leu Ser Gly Met Leu
 340 345 350
 Cys Ala Gly Val Pro Gln Gly Gly Val Asp Ala Cys Gln Gly Asp Ser
 355 360 365
 Gly Gly Pro Leu Val Gln Glu Asp Ser Arg Arg Leu Trp Phe Ile Val
 370 375 380
 Gly Ile Val Ser Trp Gly Asp Gln Cys Gly Leu Pro Asp Lys Pro Gly
 385 390 395 400
 Val Tyr Thr Arg Val Thr Ala Tyr Leu Asp Trp Ile Arg Gln Gln Thr
 405 410 415

Gly Ile

<210> 84

<211> 489

<212> DNA

<213> Homo sapiens

<400> 84

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aattacagag gagtttcatg gcactaagtc aagatattca gaaaacaata aagaagacag 180
cacgtcgga gcagcttatg agagaagaag ctgaacagaa acgtttaaaa actgtacttg 240
agctacagta tgttttggac aaattgggag atgatgaagt gcggactgac ctgaaacaag 300
gtttgaatgg agtgccaata ttgtccgaag aggagtgtg attgttggat gaattctata 360
agctagtaga ccctgaacgg gacatgagct tgaggttgaa tgaacagtat gaacatgcct 420
ccattcacct gtgggacctg ctggaaggga aggaaaaacc tgtatgtgga accacctata 480
aagttctaa
```

489

<210> 85

<211> 304

<212> DNA

<213> Homo sapiens

<400> 85

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acgcggacag cgtggccgag ctgggggagc agatcgacaa cctgcagcgg gtgaagcaga 120
agctggagaa ggagaagagc gagatgaaga tggagatcga tgacctcgct tgtaacatgg 180
aggtcatctc caaatctaag ggaaaccttg agaagatgtg ccgcacactg gaggaccaag 240
tgagtgaagt gaagaccagc gaggaggaac agcagcggct gatcaatgaa ctgactgcgc 300
agag
```

304

<210> 86

<211> 296

<212> DNA

<213> Homo sapiens

<400> 86

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ttccttaagg attaaaatgt ttagggaac acgtgttact tccacttcca gatttctgaa 120
tccatagtgt gtatgtttcc ttgtctccc aggggttgtg atcctggcag tccccatagc 180
tctacttggt tactttttag cttttgatca aaaatcttac ttttattgga gcaattttcc 240
actcccaaat gttgaatata atagtccgtt taattcccc gtttcaccgg gaattc 296
```

<210> 87

<211> 904

<212> DNA

<213> Homo sapiens

<400> 87

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tatctgatcg ttctaaaaaa gagttgtccc cggttttaac cagtgaagtt catagtgttc 180
gtgcaggacg gcatcttgct accaaattga atattttagt acagcaacat tttgacttgg 240
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attatgatgt ggagctactt catcacaag atgcacatgt agatttcctg aaaagtggtg 360
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attcgcatct aggtggcggc agtcgagaag gctcgtttta agaaacaata acattaaagt 420
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 ttggtgaatt tatgagggga aaacagatta actccttttc tacaccccag atataaaatc 780
 gatggaagtc ttgaggtccc tttggaaccg agccaaaaga tcagttaaaa aaacataccc 840
 gttactggcc tatgatttca aaaaccacc atttttaaca tgcaagcggg agttccgtta 900
 acca 904

<210> 88

<211> 387

<212> DNA

<213> Homo sapiens

<400> 88

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 ccgctgcgat tcagaagatg ttgatgaca ataaccatct tattcagtgt ataattggact 180
 ctccagaataa aggaagacc tcagagtgtt ctccagtatca gcagatgttg cacacaaact 240
 tgggtatacct tgctacaata gcagattcta atcaaaatat gcagtctctt ttaccagcac 300
 caccacaca gaatatgcct atgggtcctg gagggatgaa tcagagcggg cctccccac 360
 ctccacgctc tcacaacatg ctttcaa 387

<210> 89

<211> 481

<212> DNA

<213> Homo sapiens

<400> 89

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 tgaaccagca agctatacag attcttgaaa agattttctca gccagtgggtg gtgggtggcca 180
 ttgtaggact gtaccgtaca gggaaatcct acttgatgaa ccatctggca ggacagaatc 240
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 tgccccaccc atccaagcca aaccacaccc tggctccttct ggacaccgaa ggtctgggcg 360
 atgtggaaaa ggggtgaccct aagaatgact cctggatctt tgccctggct gtgctcctgt 420
 gcagcacctt tgtctacaac agcatgagca ccatcaacca ccaggccctg gagcagctgc 480
 a 481

<210> 90

<211> 491

<212> DNA

<213> Homo sapiens

<400> 90

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 tggaaaaggg tgaccctaag aatgactcct ggatctttgc cctggctgtg ctctgtgca 420
 gcacctttgt ctacaacagc atgagcacca tcaaccacca agccctggag cagctgcatt 480

atgtgacgga c

491

<210> 91

<211> 488

<212> DNA

<213> Homo sapiens

<400> 91

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tggggaaggt gaaggtcggg gtcaacggat ttggctcgat tgggcgcctg gtcaccaggg 120
ctgcttttaa ctctggtaaa gtggatattg ttgccatcaa tgaccccttc attgacctca 180
actacatggt ttacatgttc caatatgatt ccacccatgg caaattccat ggcaccgtcg 240
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tctctgcccc tctgctgatg ccccatgttc gtcatgggtg tgaaccatga gaagtatgac 480
acagcctc                                     488

```

<210> 92

<211> 384

<212> DNA

<213> Homo sapiens

<400> 92

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gacagtcagc cgcattctct tttgcgtcgc cagccgagcc acatcgctca gacaccatgg 60
ggaggtgtaa ggtcggagtc aacggatttg gtcgtattgg gcgcctgggc accagggctg 120
cttttaactc tggtaaagtg gatattgttg ccatcaatga ccccttcatt gacctcaact 180
acatgggtta catgttccaa tatgattcca cccatggcaa attccatggc accgtcgagg 240
ctgagaacgg gaagcttgct atcaatggaa atcccatcac catcttcag gagcgagatc 300
cctccaaaat caagtggggc gatactggcg ctgagtacgt cgtggagtcc actggcgctc 360
tcaccaccat ggagaaggct gggg                                     384

```

<210> 93

<211> 162

<212> PRT

<213> Homo sapiens

<400> 93

```

Lys Gly Lys Leu Asp Asp Tyr Gln Glu Arg Met Asn Lys Gly Glu Arg
 1           5           10           15
Leu Asn Gln Asp Gln Leu Asp Ala Val Ser Lys Tyr Gln Glu Val Thr
          20           25           30
Asn Asn Leu Glu Phe Ala Lys Glu Leu Gln Arg Ser Phe Met Ala Leu
          35           40           45
Ser Gln Asp Ile Gln Lys Thr Ile Lys Lys Thr Ala Arg Arg Glu Gln
          50           55           60
Leu Met Arg Glu Glu Ala Glu Gln Lys Arg Leu Lys Thr Val Leu Glu
          65           70           75           80
Leu Gln Tyr Val Leu Asp Lys Leu Gly Asp Asp Glu Val Arg Thr Asp
          85           90           95

```

Leu Lys Gln Gly Leu Asn Gly Val Pro Ile Leu Ser Glu Glu Glu Leu
 100 105 110

Ser Leu Leu Asp Glu Phe Tyr Lys Leu Val Asp Pro Glu Arg Asp Met
 115 120 125

Ser Leu Arg Leu Asn Glu Gln Tyr Glu His Ala Ser Ile His Leu Trp
 130 135 140

Asp Leu Leu Glu Gly Lys Glu Lys Pro Val Cys Gly Thr Thr Tyr Lys
 145 150 155 160

Val Leu

<210> 94

<211> 100

<212> PRT

<213> Homo sapiens

<400> 94

Asp Leu Glu Glu Ala Thr Leu Gln His Glu Ala Thr Ala Ala Thr Leu
 1 5 10 15

Arg Lys Lys His Ala Asp Ser Val Ala Glu Leu Gly Glu Gln Ile Asp
 20 25 30

Asn Leu Gln Arg Val Lys Gln Lys Leu Glu Lys Glu Lys Ser Glu Met
 35 40 45

~~Lys Met Glu Ile Asp Asp Leu Ala Cys Asn Met Glu Val Ile Ser Lys~~
 50 55 60

Ser Lys Gly Asn Leu Glu Lys Met Cys Arg Thr Leu Glu Asp Gln Val
 65 70 75 80

Ser Glu Leu Lys Thr Gln Glu Glu Glu Gln Gln Arg Leu Ile Asn Glu
 85 90 95

Leu Thr Ala Gln
 100

<210> 95

<211> 99

<212> PRT

<213> Homo sapiens

<400> 95

Lys Ile Leu Pro Leu Asn Gly Asn Leu Gln Ala Val Glu Leu Gly Glu
 1 5 10 15

Lys Arg Thr Ser Ser Leu Arg Ile Lys Met Phe Arg Ala Thr Arg Val
 20 25 30

Thr Ser Thr Ser Arg Phe Leu Asn Pro Tyr Val Val Cys Phe Leu Val
 35 40 45
 Leu Pro Gly Val Val Ile Leu Ala Val Pro Ile Ala Leu Leu Val Tyr
 50 55 60
 Phe Leu Ala Phe Asp Gln Lys Ser Tyr Phe Tyr Trp Ser Asn Phe Pro
 65 70 75 80
 Leu Pro Asn Val Glu Tyr Asn Ser Pro Phe Asn Ser Pro Ala Ser Pro
 85 90 95
 Gly Ile Pro

<210> 96
 <211> 257
 <212> PRT
 <213> Homo sapiens

<400> 96
 Val Gln Glu Thr Ile His Glu His Asn Lys Leu Ala Ala Asn Ser Asp
 1 5 10 15
 His Leu Met Gln Ile Gln Lys Cys Glu Leu Val Leu Ile His Thr Tyr
 20 25 30
 Pro Val Gly Glu Asp Ser Leu Val Ser Asp Arg Ser Lys Lys Glu Leu
 35 40 45
 Ser Pro Val Leu Thr Ser Glu Val His Ser Val Arg Ala Gly Arg His
 50 55 60
 Leu Ala Thr Lys Leu Asn Ile Leu Val Gln Gln His Phe Asp Leu Ala
 65 70 75 80
 Ser Thr Thr Ile Thr Asn Ile Pro Met Lys Glu Glu Gln His Ala Asn
 85 90 95
 Thr Ser Ala Asn Tyr Asp Val Glu Leu Leu His His Lys Asp Ala His
 100 105 110
 Val Asp Phe Leu Lys Ser Gly Asp Ser His Leu Gly Gly Gly Ser Arg
 115 120 125
 Glu Gly Ser Phe Lys Glu Thr Ile Thr Leu Lys Trp Cys Thr Pro Arg
 130 135 140
 Thr Asn Asn Ile Glu Leu His Tyr Cys Thr Gly Ala Tyr Arg Ile Ser
 145 150 155 160
 Pro Val Asp Val Asn Ser Arg Pro Ser Ser Cys Leu Thr Asn Phe Leu
 165 170 175

Leu Asn Gly Arg Ser Val Leu Leu Glu Gln Pro Arg Lys Ser Gly Ser
 180 185 190
 Lys Val Ile Ser His Met Leu Ser Ser His Gly Gly Glu Ile Phe Leu
 195 200 205
 His Val Leu Ser Ser Ser Arg Ser Ile Leu Glu Asp Pro Pro Ser Ile
 210 215 220
 Ser Glu Gly Cys Gly Gly Arg Val Thr Asp Tyr Arg Ile Thr Asp Phe
 225 230 235 240
 Gly Glu Phe Met Arg Gly Lys Gln Ile Asn Ser Phe Ser Thr Pro Gln
 245 250 255
 Ile

<210> 97
 <211> 128
 <212> PRT
 <213> Homo sapiens

<400> 97
 Ser Leu Pro Gln Phe Ala Val His Pro Glu Arg Ser Gly Leu Ala Asp
 1 5 10 15
 Ser Gly Asp Gly Gly Asn Met Ser Val Ala Phe Ala Ala Pro Arg Gln
 20 25 30
 Arg Gly Lys Gly Glu Ile Thr Pro Ala Ala Ile Gln Lys Met Leu Asp
 35 40 45
 Asp Asn Asn His Leu Ile Gln Cys Ile Met Asp Ser Gln Asn Lys Gly
 50 55 60
 Lys Thr Ser Glu Cys Ser Gln Tyr Gln Gln Met Leu His Thr Asn Leu
 65 70 75 80
 Val Tyr Leu Ala Thr Ile Ala Asp Ser Asn Gln Asn Met Gln Ser Leu
 85 90 95
 Leu Pro Ala Pro Pro Thr Gln Asn Met Pro Met Gly Pro Gly Gly Met
 100 105 110
 Asn Gln Ser Gly Pro Pro Pro Pro Pro Arg Ser His Asn Met Pro Ser
 115 120 125

<210> 98
 <211> 159
 <212> PRT
 <213> Homo sapiens

<400> 98

Phe Leu Asp Leu Arg Cys Tyr Arg Ala Gly Ser Ser Arg Leu Ala Val
 1 5 10 15
 Ala Met Glu Ser Gly Pro Lys Met Leu Ala Pro Val Cys Leu Val Glu
 20 25 30
 Asn Asn Asn Glu Gln Leu Leu Val Asn Gln Gln Ala Ile Gln Ile Leu
 35 40 45
 Glu Lys Ile Ser Gln Pro Val Val Val Val Ala Ile Val Gly Leu Tyr
 50 55 60
 Arg Thr Gly Lys Ser Tyr Leu Met Asn His Leu Ala Gly Gln Asn His
 65 70 75 80
 Gly Phe Pro Leu Gly Ser Thr Val Gln Ser Glu Thr Lys Gly Ile Trp
 85 90 95
 Met Trp Cys Val Pro His Pro Ser Lys Pro Asn His Thr Leu Val Leu
 100 105 110
 Leu Asp Thr Glu Gly Leu Gly Asp Val Glu Lys Gly Asp Pro Lys Asn
 115 120 125
 Asp Ser Trp Ile Phe Ala Leu Ala Val Leu Leu Cys Ser Thr Phe Val
 130 135 140
 Tyr Asn Ser Met Ser Thr Ile Asn His Gln Ala Leu Glu Gln Leu
 145 150 155

<210> 99

<211> 147

<212> PRT

<213> Homo sapiens

<400> 99

Met Glu Ser Gly Pro Lys Met Leu Ala Pro Val Cys Leu Val Glu Asn
 1 5 10 15
 Asn Asn Glu Gln Leu Leu Val Asn Gln Gln Ala Ile Gln Ile Leu Glu
 20 25 30
 Lys Ile Ser Gln Pro Val Val Val Val Ala Ile Val Gly Leu Tyr Arg
 35 40 45
 Thr Gly Lys Ser Tyr Leu Met Asn His Leu Ala Gly Gln Asn His Gly
 50 55 60
 Phe Pro Leu Gly Ser Thr Val Gln Ser Glu Thr Lys Gly Ile Trp Met
 65 70 75 80
 Trp Cys Val Pro His Pro Ser Lys Pro Asn His Thr Leu Val Leu Leu

85 90 95
 Asp Thr Glu Gly Leu Gly Asp Val Glu Lys Gly Asp Pro Lys Asn Asp
 100 105 110
 Ser Trp Ile Phe Ala Leu Ala Val Leu Leu Cys Ser Thr Phe Val Tyr
 115 120 125
 Asn Ser Met Ser Thr Ile Asn His Gln Ala Leu Glu Gln Leu His Tyr
 130 135 140
 Val Thr Asp
 145

<210> 100
 <211> 124
 <212> PRT
 <213> Homo sapiens

<400> 100
 Met Gly Lys Val Lys Val Gly Val Asn Gly Phe Gly Arg Ile Gly Arg
 1 5 10 15
 Leu Val Thr Arg Ala Ala Phe Asn Ser Gly Lys Val Asp Ile Val Ala
 20 25 30
 Ile Asn Asp Pro Phe Ile Asp Leu Asn Tyr Met Val Tyr Met Phe Gln
 35 40 45
 Tyr Asp Ser Thr His Gly Lys Phe His Gly Thr Val Glu Ala Glu Asn
 50 55 60
 Gly Lys Leu Val Ile Asn Gly Asn Pro Ile Thr Ile Phe Gln Glu Arg
 65 70 75 80
 Asp Pro Ser Lys Ile Lys Trp Gly Asp Ala Gly Ala Glu Tyr Val Val
 85 90 95
 Glu Ser Thr Gly Val Phe Thr Thr Met Glu Lys Ala Gly Ala His Leu
 100 105 110
 Gln Gly Gly Ala Lys Arg Val Ile Ile Ser Ala Pro
 115 120

<210> 101
 <211> 127
 <212> PRT
 <213> Homo sapiens

<400> 101
 Gln Ser Ala Ala Ser Ser Phe Ala Ser Pro Ala Glu Pro His Arg Ser
 1 5 10 15

Asp Thr Met Gly Lys Val Lys Val Gly Val Asn Gly Phe Gly Arg Ile
20 25 30

Gly Arg Leu Val Thr Arg Ala Ala Phe Asn Ser Gly Lys Val Asp Ile
35 40 45

Val Ala Ile Asn Asp Pro Phe Ile Asp Leu Asn Tyr Met Val Tyr Met
50 55 60

Phe Gln Tyr Asp Ser Thr His Gly Lys Phe His Gly Thr Val Glu Ala
65 70 75 80

Glu Asn Gly Lys Leu Val Ile Asn Gly Asn Pro Ile Thr Ile Phe Gln
85 90 95

Glu Arg Asp Pro Ser Lys Ile Lys Trp Gly Asp Thr Gly Ala Glu Tyr
100 105 110

Val Val Glu Ser Thr Gly Val Phe Thr Thr Met Glu Lys Ala Gly
115 120 125

<210> 102

<211> 1225

<212> DNA

<213> Homo sapiens

<400> 102

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atggcggcgc ggtcgtcgtc gggggtggcg gcggcagagg gggcggcggc cctggcggca 60
gcggagacgg cagccgtgac ggtggcagcg gcggcggcgg acctgggcct gggggaatga 120
ggcggccgcg gcgggcccagc ggaggagccg tgtagcggag aagctcccc tccctgcttc 180
ccttgccga gccgggggcg cgcgcgcacg cggccgtcca gagcgggctc cccacccctc 240
gactcctgcg acccgccacc cacccccacc cggggccgga ggatgatgaa gctcaagtcg 300
aaccagaccc gcacctacga cggcgacggc tacaagaagc gggccgcatg cctgtgtttc 360
cgcagcgaga gcgaggagga ggtgctactc gtgagcagta gtcgccatcc agacagatgg 420
attgtccctg gaggaggcat ggagcccagc gaggagccaa gtgtggcagc agttcgtgaa 480
gtctgtgagg aggctggagt aaaagggaca ttgggaagat tagttggaat ttttgagaac 540
caggagagga agcacaggac gtatgtctat gtgtcattg tcactgaagt gctggaagac 600
tggaagatt cagttaacat tggaaggaag agggaatggt taaaataga agacgccata 660
aaagtgtgc agtatcaca acccgtgcag gcatcatatt ttgaaacatt gaggcaaggc 720
tactcagcca acaatggcac cccagtcgtg gccaccacat actcggtttc tgctcagagc 780
tcgatgtcag gcatcagatg actgaagact tcctgtaaga gaaatggaaa ttggaaacta 840
gactgaagtg caaatcttcc ctctcaccct ggctctttcc acttctcaca ggcctcctct 900
ttcaaataag gcatggtggg cagcaaagaa aggggtgtatt gataatgttg ctgtttggtg 960
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aggccaacag ccttcccctg ccttggattc tgaagtgttc ctgtttgtct tatcctggcc 1140
ctggccagac gttttctttg atttttaatt tttttttttt attaaaagat accagtatga 1200
gaaaaaaaaa aaaaaaaaaa tcgag 1225

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<210> 103

<211> 741

<212> DNA

<213> Homo sapiens

<400> 103

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agaaacctca atcggattca gcaaaggaat ggtgttatta tcactacata ccaaattgtta 60
atcaataact ggcagcaact ttcaagcttt aggggccaaag agtttgtgtg ggactatgtc 120
atcctcgatg aagcacataa aataaaaacc tcactacta agtcagcaat atgtgtcgt 180
gctattcctg caagtaatcg cctcctcctc acaggaaccc caatccagaa taatttacia 240
gaactatggt ccctatttga ttttgcttgt caagggtccc tgctgggaac attaaaaact 300
tttaagatgg agtatgaaaa tcctattact agagcaagag agaaggatgc taccacagga 360
gaaaaagcct tgggatttaa aatatctgaa aacttaatgg caatcataaa accctatttt 420
ctcaggagga ctaaaagaaga cgtacagaag aaaaagtcaa gcaaccaga ggccagactt 480
aatgaaaaga atccagatgt tgatgccatt tgtgaaatgc cttccctttc caggagaaat 540
gatttaatta tttggatacg acttgtgcct ttacaagaag aaatatacag gaaatttgtg 600
tctttagatc atatcaagga gttgctaag gagacgcgt cacttttggc tgagctaggt 660
gtcttaaga agctgtgtga tcactcctagg ctgctgtctg cacgggcttg ttgtttgcta 720
aatcttgagg cattctctgc t 741

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<210> 104

<211> 321

<212> DNA

<213> Homo sapiens

<400> 104

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ttgtctcg tcatcaaaga caccaaactg ctgtgctata aaagttccaa ggaccagcag 60
cctcagatgg aactgccact ccaaggctgt aacattacgt acatcccga agacagcaaa 120
aagaagaagc acgagctgaa gattactcag cagggcacgg acccgcttgt tctcgccgtc 180
cagagcaagg aacaggccga gcagtggctg aaggatgatc aagaagccta cagtgggtgt 240
agtggccccc tggattcaga gtgtcctcct ccaccaagct ccccggtgca caaggcagaa 300
ctggagaaga aactgtcttc a 321

```

<210> 105

<211> 389

<212> DNA

<213> Homo sapiens

<400> 105

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cagcactggc cacactataa aattcaggtt cagaaaaaca gggtaagtca cagacagcaa 60
cgcttccagc atttattttc tttgcaccca tgggcaattt gagaaaattt accttttagaa 120
cgaactctgt taaagggtaca gacagtacaa tactttttat tcagaagggt tctgcataaa 180
gggtgatagtc ttttgactta atatattatt gtctcctgcc ttgtgtttct ggaatgaatg 240
aaggtcatta tttagaagat aatctgggtt gtatttgtgt cgtcagattg aattttcatt 300
gcacatgcta cttaatgtct ttaccaata ataacaaagg gaaagaaaac caaatataga 360
tgtataataa ggaaaagctg gcctataga 389

```

<210> 106

<211> 446

<212> DNA

<213> Homo sapiens

<400> 106

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gccacatttg ccctgggtcat agtttaaaaca ccaggctctg tgtcacatct ttttgggtgcc 60
acaagtatca ctccattggt cagagagtaa tgtattagtt ctgcccattt cattcttcac 120
ttttatttct tccatttcat tagcatttat atcagctcaa gaagttaagg ttagaaaatt 180
ttccacttca aattttcagt acagaaatgt gctgtgatgt ttgacaagac tatttcatag 240
taagtgaagt aatgtttatt ggcctctgct ctctctgtg tcagacctag gaagcctgag 300
gattacttag ttgttctgtc tctgggtcca caggcagaat ttggcccatc caaagactgg 360
ccaagtgcc aaaaaaggcc tgattaggcc ctgaaattca gtgaaattct gcctgaagaa 420

```

acctcttatt gaatttgaaa accata:

446

<210> 107

<211> 467

<212> DNA

<213> Homo sapiens

<400> 107

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ccgccgctgc cgtcgcccttc ctgggattgg agtctcgagc tttcttcggt cggtcgccgg 60
ggggttcgcg cccttctcgc gcctcggggc tgcgaggctg ggggaagggg tggagggggc 120
tggtgatcgc cgcgtttaag ttgcgctcgg ggcggccatg tcggccggcg aggtcgagcg 180
cctagtgtcg gagctgagcg gcgggaccgg aggggatgag gaggaagagt ggctctatgg 240
cgatgaagat gaagttgaaa ggcagaaga agaaaatgcc agtgctaata ctccatctgg 300
aattgaagat gaaactgctg aaaatgggtg accaaaaccg aaagtgactg agaccgaaga 360
tgatagtgat agtgacagcg atgatgatga agatgatgtg catgtcacta taggagacat 420
taaaacggga gcaccacagt atgggagtta tggtagagca cctgtaa 467

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<210> 108

<211> 491

<212> DNA

<213> Homo sapiens

<400> 108

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gaaagataca acttcccca cccaaaccgg tttgtggagg acgacatgga taagaatgaa 60
atcgccctctg ttgcgtaccg ttaccgcagg tggagcttg gagatgatat tgaccttatt 120
gtccgttgtg agcacgatgg cgtcatgact ggagccaacg gggaaagtgtc ctccatcaac 180
atcaagacac tcaatgagtg ggattccagg cactgtaatg gcgttgactg gcgtcagaag 240
ctggactctc agcgagggggc tgtcattgcc acggagctga agaacaacag ctacaagttg 300
gcccgggtgga cctgctgtgc tttgtgtggc ggatctgagt acctcaagct tgggtatgtg 360
tctcggtacc acgtgaaaga ctctcacgc cagtcaccc taggcacca gcagttcaag 420
cctaattgagt ttgccagcca gatcaacctg agcgtggaga atgcctgagg cattttacgc 480
tgcgtcattg a 491

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<210> 109

<211> 489

<212> DNA

<213> Homo sapiens

<400> 109

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ctcagatagt actgaaccct ttatcaacta tgttttttca gtctgacaac caaggcggct 60
actaagtgac taaggggagc gtagtataca gtgtggataa gcaggacaaa ggggtgattc 120
acatcccagg caggacagag caggagatca tgagatttca tcactcagga tggcttgatg 180
tttattttat tttattcttt ttttttttg agatggagtc tcactcttgc ccaggctgga 240
gtgcagtggg gcgatcttgg ctactgcaa cctctgcctc ctgggttcaa gcagttctcc 300
tgctcagcc tccaagtag ctgggattac aggcgtccgc caccatgcc agccaatttt 360
tgtactttta gtagagatgg ggtttcacca tgttggccag gctgggtctg aactcctgac 420
ctcaggtgat ccactgcct cggcctccca aagtgtctgg attataggca tgcgccacca 480
tgcccgggc 489

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<210> 110

<211> 391

<212> DNA

<213> Homo sapiens

<400> 110

gcggagtcog ctggctgacc cgagcgctgg tctccgccgg gaaccctggg gcatggagag 60
 gtctgagtag ctccggcccg gcgcacgctg catcgccggag ccaggctgcc gctgtcccag 120
 tggagttcca ggagcaccac ctgagtgagg tgcagaatat ggcattctgag gagaagctgg 180
 agcaggtgct gagttccatg aaggagaaca aagtggccat cattggaaag attcataccc 240
 cgatggagta taagggggag ctagcctcct atgatatgcg gctgaggcgt aagtggact 300
 tatttgccaa cgtaatccat gtgaagtcac ttcctgggta tatgactcgg cacaacaatc 360
 tagacctggg gatcattcga gagcagacag a 391

<210> 111

<211> 172

<212> PRT

<213> Homo sapiens

<400> 111

Met Met Lys Leu Lys Ser Asn Gln Thr Arg Thr Tyr Asp Gly Asp Gly
 1 5 10 15

Tyr Lys Lys Arg Ala Ala Cys Leu Cys Phe Arg Ser Glu Ser Glu Glu
 20 25 30

Glu Val Leu Leu Val Ser Ser Ser Arg His Pro Asp Arg Trp Ile Val
 35 40 45

Pro Gly Gly Gly Met Glu Pro Glu Glu Glu Pro Ser Val Ala Ala Val
 50 55 60

Arg Glu Val Cys Glu Glu Ala Gly Val Lys Gly Thr Leu Gly Arg Leu
 65 70 75 80

Val Gly Ile Phe Glu Asn Gln Glu Arg Lys His Arg Thr Tyr Val Tyr
 85 90 95

Val Leu Ile Val Thr Glu Val Leu Glu Asp Trp Glu Asp Ser Val Asn
 100 105 110

Ile Gly Arg Lys Arg Glu Trp Phe Lys Ile Glu Asp Ala Ile Lys Val
 115 120 125

Leu Gln Tyr His Lys Pro Val Gln Ala Ser Tyr Phe Glu Thr Leu Arg
 130 135 140

Gln Gly Tyr Ser Ala Asn Asn Gly Thr Pro Val Val Ala Thr Thr Tyr
 145 150 155 160

Ser Val Ser Ala Gln Ser Ser Met Ser Gly Ile Arg
 165 170

<210> 112

<211> 247

<212> PRT

<213> Homo sapiens

<400> 112

Arg Asn Leu Asn Arg Ile Gln Gln Arg Asn Gly Val Ile Ile Thr Thr


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<210> 113
<211> 107
<212> PRT
<213> Homo sapiens
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<400> 113
Leu Leu Cys Val Ile Lys Asp Thr Lys Leu Leu Cys Tyr Lys Ser Ser

1 5 10 15
 Lys Asp Gln Gln Pro Gln Met Glu Leu Pro Leu Gln Gly Cys Asn Ile
 20 25 30
 Thr Tyr Ile Pro Lys Asp Ser Lys Lys Lys Lys His Glu Leu Lys Ile
 35 40 45
 Thr Gln Gln Gly Thr Asp Pro Leu Val Leu Ala Val Gln Ser Lys Glu
 50 55 60
 Gln Ala Glu Gln Trp Leu Lys Val Ile Lys Glu Ala Tyr Ser Gly Cys
 65 70 75 80
 Ser Gly Pro Val Asp Ser Glu Cys Pro Pro Pro Pro Ser Ser Pro Val
 85 90 95
 His Lys Ala Glu Leu Glu Lys Lys Leu Ser Ser
 100 105

 <210> 114
 <211> 155
 <212> PRT
 <213> Homo sapiens

 <400> 114
 Glu Arg Tyr Asn Phe Pro Asn Pro Asn Pro Phe Val Glu Asp Asp Met
 1 5 10 15
 Asp Lys Asn Glu Ile Ala Ser Val Ala Tyr Arg Tyr Arg Arg Trp Lys
 20 25 30
 Leu Gly Asp Asp Ile Asp Leu Ile Val Arg Cys Glu His Asp Gly Val
 35 40 45
 Met Thr Gly Ala Asn Gly Glu Val Ser Phe Ile Asn Ile Lys Thr Leu
 50 55 60
 Asn Glu Trp Asp Ser Arg His Cys Asn Gly Val Asp Trp Arg Gln Lys
 65 70 75 80
 Leu Asp Ser Gln Arg Gly Ala Val Ile Ala Thr Glu Leu Lys Asn Asn
 85 90 95
 Ser Tyr Lys Leu Ala Arg Trp Thr Cys Cys Ala Leu Leu Ala Gly Ser
 100 105 110
 Glu Tyr Leu Lys Leu Gly Tyr Val Ser Arg Tyr His Val Lys Asp Ser
 115 120 125
 Ser Arg His Val Ile Leu Gly Thr Gln Gln Phe Lys Pro Asn Glu Phe
 130 135 140
 Ala Ser Gln Ile Asn Leu Ser Val Glu Asn Ala

145

150

155

<210> 115

<211> 129

<212> PRT

<213> Homo sapiens

<400> 115

Gly Val Arg Trp Leu Thr Arg Ala Leu Val Ser Ala Gly Asn Pro Gly
 1 5 10 15

Ala Trp Arg Gly Leu Ser Thr Ser Ala Ala Ala His Ala Ala Ser Arg
 20 25 30

Ser Gln Ala Ala Ala Val Pro Val Glu Phe Gln Glu His His Leu Ser
 35 40 45

Glu Val Gln Asn Met Ala Ser Glu Glu Lys Leu Glu Gln Val Leu Ser
 50 55 60

Ser Met Lys Glu Asn Lys Val Ala Ile Ile Gly Lys Ile His Thr Pro
 65 70 75 80

Met Glu Tyr Lys Gly Glu Leu Ala Ser Tyr Asp Met Arg Leu Arg Arg
 85 90 95

Lys Leu Asp Leu Phe Ala Asn Val Ile His Val Lys Ser Leu Pro Gly
 100 105 110

Tyr Met Thr Arg His Asn Asn Leu Asp Leu Val Ile Ile Arg Glu Gln
 115 120 125

Thr

<210> 116

<211> 550

<212> DNA

<213> Homo sapiens

<400> 116

gaattcggca ccagcctcag agccccccag cccgggtacc accccctgcg gaaaggtacc 60
 catctgcatt cctgcccgtc gggacctggt ggacagteca gcctccttgg cctctagcct 120
 tggctcaccg ctgacctagag ccaaggagct catcctgaat gaccttcccg ccagcactcc 180
 tgcttcaaaa tctgtgact cctccccgcc ccaggacgct tccaccccca ggcccagctc 240
 ggccagtcac ctctgccagc ttgctgccaa gccagcacct tccacggaca gcgtcgccct 300
 gaggagcccc ctgactctgt ccagtcctct caccacgtcc ttcagcctgg gctccacacag 360
 cactctcaac ggagacctct ccgtgcccag ctctacgtc agcctccacc tgtcccccca 420
 ggtcagcagc tctgtggtgt acggacgtc ccccgatgat gcatttgagt ctcatcccca 480
 tctccgaggg tcatccgtct ctctctccct acccagcatc cctgggggaa agccggccta 540
 ctcttccac 550

<210> 117

<211> 154

<212> DNA

<213> Homo sapiens

<400> 117

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ttctgagggga aagccgagtg gagtgggcca cccggcgggcg gtgacaatga gttttcttgg 60
aggctttttt ggtccattt gtgagattga tgttgccctt aatgatgggg aaaccaggaa 120
aatggcagaa atgaaaactg aggatggcaa agta 154

```

<210> 118

<211> 449

<212> DNA

<213> Homo sapiens

<400> 118

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gaattcggca ccagggcccgc cagcccagtg gtcgccgccca tggcttcgcc gcagctctgc 60
cgcgcgctgg tgctggcgca atgggtggcg gaggcgctgc gggccccgcg cgctgggcag 120
cctctgcagc tgctggacgc ctcttggtac ctgccgaagc tggggcgcca cgcgcgacgc 180
gagttcgagg agcgccacat cccggcgccc gctttcttcg acatcgacca gtgcagcgac 240
cgcacctcgc cctacgacca catgctgccc gggggcgagc atttcgcgga gtacgcaggc 300
cgcttggggc tggggcgggc caccacgctc gtgatctacg acgccagcga ccagggcctc 360
tactccgccc cgcgcgctcg gtggatgttc cgcgccctcg gccaccacgc cgtgtcactg 420
cttgatggcg gcctccgccca ctggctgcg 449

```

<210> 119

<211> 642

<212> DNA

<213> Homo sapiens

<400> 119

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gaattcggca cgagcagtaa cccgaccgcc gctgggtcttc gctggacacc atgaatcaca 60
ctgtccaaac cttcttctct cctgtcaaca gtggccagcc ccccaactat gagatgctca 120
aggaggagca cgaggtggct gtgctggggg cgcccacaaa ccctgctccc cggacgtcca 180
ccgtgatcca catccgcagc gagacctccg tgcccagcca tgctgtctgg tccctgttca 240
acacctctt catgaacccc tgctgcctgg gcttcatagc attcgcctac tccgtgaagt 300
ctagggacag gaagatggtt ggccgactga ccggggccca ggccatagcc tccaccgcca 360
agtgcctgaa catctgggccc ctgattctgg gcattctcat gaccattctg ctcatcgta 420
tcccagtgct gatcttccag gcctatggat agatcaggag gcatcactga ggccaggagc 480
tctgcccatt acctgtatcc cacgtactcc aacttccatt cctcgccctg ccccgaggc 540
cgagtcctgt atcagccctt tatcttcaca cgcttttcta caatggcatt caataaagtg 600
cacgtgtttc tggtgaaaaa aaaaaaaaaa aaaaaactcg ag 642

```

<210> 120

<211> 603

<212> DNA

<213> Homo sapiens

<400> 120

```

gaattcggca cgagccacaa cagccactac gactgcatcc actggatcca cggccacccc 60
gtcctccacc ccgggaacag ctccccctcc caaagtgtcg accagcccgg ccaccacacc 120
catgtccacc atgtccacaa tccacacctc ctctactcca gagaccaccc acacctccac 180
agtgtgacc accacagcca ccatgacaag ggccaccaat tccacggcca caccctcctc 240
cactctgggg acgaccgggä tctcactga cctgaccaca acagccacta caactgcagc 300
cactggatcc acggccaccc tgtcctccac cccagggacc acctggatcc tcacagagcc 360
gagcactata gccaccgtga tgggtgccac cggttccacg gccaccgcct cctccactct 420
gggaacagct cacaccccca aagtgggtgac caccatggcc actatgcccc cagccactgc 480

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ctccacgggtt cccagctcgt ccaccgtggg gaccaccgc acccctgcag tgctccccag 540
 cagcctgccca accttcagcg tgtccactgt gtcctcctca gtcctcacca ccctgagacc 600
 cac 603

<210> 121
 <211> 178
 <212> PRT
 <213> Homo sapiens

<400> 121
 Ser Glu Pro Pro Ser Pro Ala Thr Thr Pro Cys Gly Lys Val Pro Ile
 1 5 10 15
 Cys Ile Pro Ala Arg Arg Asp Leu Val Asp Ser Pro Ala Ser Leu Ala
 20 25 30
 Ser Ser Leu Gly Ser Pro Leu Pro Arg Ala Lys Glu Leu Ile Leu Asn
 35 40 45
 Asp Leu Pro Ala Ser Thr Pro Ala Ser Lys Ser Cys Asp Ser Ser Pro
 50 55 60
 Pro Gln Asp Ala Ser Thr Pro Arg Pro Ser Ser Ala Ser His Leu Cys
 65 70 75 80
 Gln Leu Ala Ala Lys Pro Ala Pro Ser Thr Asp Ser Val Ala Leu Arg
 85 90 95
 Ser Pro Leu Thr Leu Ser Ser Pro Phe Thr Thr Ser Phe Ser Leu Gly
 100 105 110
 Ser His Ser Thr Leu Asn Gly Asp Leu Ser Val Pro Ser Ser Tyr Val
 115 120 125
 Ser Leu His Leu Ser Pro Gln Val Ser Ser Ser Val Val Tyr Gly Arg
 130 135 140
 Ser Pro Val Met Ala Phe Glu Ser His Pro His Leu Arg Gly Ser Ser
 145 150 155 160
 Val Ser Ser Ser Leu Pro Ser Ile Pro Gly Gly Lys Pro Ala Tyr Ser
 165 170 175
 Phe His

<210> 122
 <211> 36
 <212> PRT
 <213> Homo sapiens

<400> 122
 Met Ser Phe Leu Gly Gly Phe Phe Gly Pro Ile Cys Glu Ile Asp Val
 1 5 10 15

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<210> 123
<211> 136
<212> PRT
<213> Homo sapiens
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<400> 123
Met Ala Ser Pro Gln Leu Cys Arg Ala Leu Val Ser Ala Gln Trp Val
1 5 10 15

Ala Glu Ala Leu Arg Ala Pro Arg Ala Gly Gln Pro Leu Gln Leu Leu
20 25 30

Asp Ala Ser Trp Tyr Leu Pro Lys Leu Gly Arg Asp Ala Arg Arg Glu
35 40 45

Phe Glu Glu Arg His Ile Pro Gly Ala Ala Phe Phe Asp Ile Asp Gln
50 55 60

Cys Ser Asp Arg Thr Ser Pro Tyr Asp His Met Leu Pro Gly Ala Glu
65 70 75 80

His Phe Ala Glu Tyr Ala Gly Arg Leu Gly Val Gly Ala Ala Thr His
85 90 95

Val Val Ile Tyr Asp Ala Ser Asp Gln Gly Leu Tyr Ser Ala Pro Arg
100 105 110

Val	Trp	Trp	Met	Phe	Arg	Ala	Phe	Gly	His	His	Ala	Val	Ser	Leu	Leu
		115					120					125			

Asp Gly Gly Leu Arg His Trp Leu
130 135

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<210> 124
<211> 133
<212> PRT
<213> Homo sapiens
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<400> 124
Met Asn His Thr Val Gln Thr Phe Phe Ser Pro Val Asn Ser Gly Gln
1 5 10 15

Pro Pro Asn Tyr Glu Met Leu Lys Glu Glu His Glu Val Ala Val Leu
20 25 30

Gly Ala Pro His Asn Pro Ala Pro Pro Thr Ser Thr Val Ile His Ile
35 40 45

Arg Ser Glu Thr Ser Val Pro Asp His Val Val Trp Ser Leu Phe Asn
 50 55 60
 Thr Leu Phe Met Asn Pro Cys Cys Leu Gly Phe Ile Ala Phe Ala Tyr
 65 70 75 80
 Ser Val Lys Ser Arg Asp Arg Lys Met Val Gly Asp Val Thr Gly Ala
 85 90 95
 Gln Ala Tyr Ala Ser Thr Ala Lys Cys Leu Asn Ile Trp Ala Leu Ile
 100 105 110
 Leu Gly Ile Leu Met Thr Ile Leu Leu Ile Val Ile Pro Val Leu Ile
 115 120 125
 Phe Gln Ala Tyr Gly
 130
 <210> 125
 <211> 195
 <212> PRT
 <213> Homo sapiens
 <400> 125
 Thr Thr Ala Thr Thr Thr Ala Ser Thr Gly Ser Thr Ala Thr Pro Ser
 1 5 10 15
 Ser Thr Pro Gly Thr Ala Pro Pro Pro Lys Val Leu Thr Ser Pro Ala
 20 25 30
 Thr Thr Pro Met Ser Thr Met Ser Thr Ile His Thr Ser Ser Thr Pro
 35 40 45
 Glu Thr Thr His Thr Ser Thr Val Leu Thr Thr Thr Ala Thr Met Thr
 50 55 60
 Arg Ala Thr Asn Ser Thr Ala Thr Pro Ser Ser Thr Leu Gly Thr Thr
 65 70 75 80
 Arg Ile Leu Thr Glu Leu Thr Thr Thr Ala Thr Thr Thr Ala Ala Thr
 85 90 95
 Gly Ser Thr Ala Thr Leu Ser Ser Thr Pro Gly Thr Thr Trp Ile Leu
 100 105 110
 Thr Glu Pro Ser Thr Ile Ala Thr Val Met Val Pro Thr Gly Ser Thr
 115 120 125
 Ala Thr Ala Ser Ser Thr Leu Gly Thr Ala His Thr Pro Lys Val Val
 130 135 140
 Thr Thr Met Ala Thr Met Pro Thr Ala Thr Ala Ser Thr Val Pro Ser
 145 150 155 160

Ser Ser Thr Val Gly Thr Thr Arg Thr Pro Ala Val Leu Pro Ser Ser
 165 170 175

Leu Pro Thr Phe Ser Val Ser Thr Val Ser Ser Ser Val Leu Thr Thr
 180 185 190

Leu Arg Pro
 195

<210> 126
 <211> 509
 <212> DNA
 <213> homo sapien

<400> 126
 gaattcggca cgagccaagt accccctgag gaatctgcag cctgcatctg agtacaccgt 60
 atccctcgtg gccataaagg gcaaccaaga gagccccaaa gccactggag tctttaccac 120
 actgcagcct gggagctcta ttccacctta caacaccgag gtgactgaga ccaccattgt 180
 gatcacatgg acgcctgctc caagaattgg ttttaagctg ggtgtacgac caagccaggg 240
 aggagaggca ccacgagaag tgacttcaga ctcaggaagc atcgttgtgt ccggcttgac 300
 tccaggagta gaatacgtct acaccatcca agtcttgaga gatggacagg aaagagatgc 360
 gccatttgta aacaaagtgg tgacaccatt gtctccacca acaaacttgc atctggaggc 420
 aaaccctgac actggagtgc tcacagtctc ctggagagga gcaccacccc agacattact 480
 gggatataga ttaccacaac ccctacaaa 509

<210> 127
 <211> 500
 <212> DNA
 <213> homo sapien

<400> 127
 gaattcggca cgagccactg atgtccgggg agtcagccag gagcttgggg aaggggaagcg 60
 cgccccggg gccgggtccc gaggggtcga tccgcatcta cagcatgagg ttctgcccgt 120
 ttgctgagag gacgcgtcta gtcctgaagg ccaagggaat caggcatgaa gtcatcaata 180
 tcaacctgaa aaataagcct gagtggttct ttaagaaaaa tccctttggt ctggtgccag 240
 ttctggaaaa cagtcagggt cagctgatct acgagtctgc catcacctgt gagtacctgg 300
 atgaagcata ccaggggaag aagctgttgc cggatgaccc ctatgagaaa gcttgccaga 360
 agatgatctt agagttgttt tctaagggtc catccttggt aggaagcttt attagaagcc 420
 aaaataaaga agactatgct ggccataaag aagaatttcg taaagaattt accaagctag 480
 aggaggttct gactaataag 500

<210> 128
 <211> 500
 <212> DNA
 <213> homo sapien

<400> 128
 agctttctct tgctgccgct cggtcacgct tgtgcccga gaggaaaca gtgacagacc 60
 tggagactgc agttctctat ccttcacaca gctctttcac catgcctgga tcacttcctt 120
 tgaatgcaga agcttgctgg ccaaaagatg tgggaattgt tgcccttgag atctattttc 180
 cttctcaata tgttgatcaa gcagagttgg aaaaatatga tgggtgagat gctggaaaagt 240
 ataccattgg cttggggccag gccaaagatg gcttctgcac agatagagaa gatattaact 300
 ctctttgcat gactgtgggt cagaatctta tggagagaaa taacctttcc tatgattgca 360

ttgggcccgt	ggaagttgga	acagagacaa	tcacgcacaa	atcaaagtct	gtgaagacta	420
atgtgatgca	gctgtttgaa	gagtcctggga	atacagatat	agaaggaatc	gacacaacta	480
atgcatgcta	tggaggcaca					500

<210> 129

<211> 497

<212> DNA

<213> homo sapien

<400> 129

gaattcggca	cgagcagagg	tctccagagc	cttctctctc	ctgtgcaaaa	tggcaactct	60
taaggaaaaa	ctcattgcac	cagttgcgga	agaagaggca	acagttccaa	acaataagat	120
cactgtagtg	ggtgttgac	aagttggtat	ggcgtgtgct	atcagcattc	tgggaaagtc	180
tctggctgat	gaacttgctc	ttgtggatgt	tttgggaagat	aagcttaaag	gagaaatgat	240
ggatctgcag	catgggagct	tatttcttca	gacacctaaa	attgtggcag	ataaagatta	300
ttctgtgacc	gccaatctca	agattgtagt	ggtaactgca	ggagtccgtc	agcaagaagg	360
ggagagtcgg	ctcaatctgg	tgcagagaaa	tgtaaatgtc	ttcaaattca	ttattcctca	420
gacgtcaag	tacagtcctg	attgcatcat	aattgtgggt	tccaaccag	tggacattct	480
tacgtatgtt	acctgga					497

<210> 130

<211> 383

<212> DNA

<213> homo sapien

<400> 130

gaattcggca	cgagggccgc	ggctgccgac	tgggtcccct	gccgctgtcg	ccaccatggc	60
tccgcaccgc	cccgcgccc	cgctgctttg	cgcgctgtcc	ctggcgctgt	gcgcgctgtc	120
gctgcccgtc	cgcgcggcca	ctgcgtcgcg	ggggcgctcc	caggcggggg	cgccccaggg	180
gcgggtgcc	gagcgcggc	ccaacagcat	ggtggtggaa	caccccgagt	tcctcaaggc	240
agggaaagag	cctggcctgc	agatctggcg	tgtggagaaa	gttcgatctg	gtggcccggtg	300
cccaccaacc	tttatggaga	cttcttcacg	ggcgacgcct	acgtcatcct	gaagacagtg	360
cagcttaaga	acggaaaatc	ttg				383

<210> 131

<211> 509

<212> DNA

<213> homo sapien

<400> 131

gaattcggca	cgagagtcag	ccgcattctc	ttttgcgtcg	ccagccgagc	cacatcgctc	60
agacaccatg	gggaagggtga	aggtcggagt	caacggattt	ggtcgtattg	ggcgccctggt	120
caccagggct	gcttttaact	ctggtaaagt	ggatattgtt	gccatcaatg	acccttcat	180
tgacctcaac	tacatggttt	acatgttcca	atatgattcc	acccatggca	aattccatgg	240
caccgtcaag	gctgagaacg	ggaagcttgt	catcaatgga	aatcccatca	ccatcttcca	300
ggagcgagat	ccctccaaaa	tcaagtgggg	cgatgctggc	gctgagtacg	tcgtggagtc	360
cactggccgt	cttcaccacc	atggagaagg	ctggggctca	tttgcagggg	ggagccaaaa	420
gggtcatcat	ctctgcccc	tctgctgacg	cccccatggt	cgtcatgggt	gtgaaccatg	480
agaagtatga	caacagcctc	aagatcatc				509

<210> 132

<211> 357

<212> DNA

<213> homo sapien

<400> 132

gaattcggca	cgagtaagaa	gaagccccta	gaccacagct	ccacaccatg	gactggacct	60
ggaggatcct	cttcttggtg	gcagcagcaa	caggtgccca	ctcccagggtg	caactgggtgc	120
aatctgggtc	tgagttgaag	aagcctgggg	cctcagtgaa	ggtttcctgc	aaggcttctg	180
gacacatctt	cagtatctat	ggtttgaatt	gggtgcgaca	ggccccctggt	caaggccttg	240
agtggatggg	atggatcaaa	gtcgacactg	cgaacccaac	gtatgccag	ggcttcacag	300
gacgatttgt	cttctccctg	gacacctctg	tcagcacggc	atatctgcag	atcagca	357

<210> 133

<211> 468

<212> DNA

<213> homo sapien

<400> 133

gaattcggca	cgaggcgccc	cgaaccgtcc	tcctgctgct	ctcggcgggc	ctggccctga	60
ccgagacctg	ggccggctcc	cactccatga	ggtatttcga	caccgccatg	tcccggccccg	120
gccgcgggga	gccccgttc	atctcagtg	gctacgtgga	cgacacgcag	ttcgtgaggt	180
tcgacagcga	cgccgcgagt	ccgagagagg	agccgcgggc	gccgtggata	gagcaggagg	240
ggccggagta	ttgggaccgg	aacacacaga	tcttcaagac	caacacacag	actgaccgag	300
agagcctgcg	gaacctgcgc	ggctactaca	accagagcga	ggccgggtct	cacaccctcc	360
agagcatgta	cggctgcgac	gtggggccgg	acgggcgcct	cctccgcggg	cataaccagt	420
acgcctacga	cggcaaggat	tacatcgccc	tgaacgagga	cctgcgct		468

<210> 134

<211> 214

<212> DNA

<213> homo sapien

<400> 134

gaattcggca	cgagctgcgt	cctgctgagc	tctgttctct	ccagcacctc	ccaaccact	60
agtgcctggt	tctcttgctc	caccaggaac	aagccaccat	gtctcgccag	tcaagtgtgt	120
ccttccggag	cgggggcagt	cgtagcttca	gcaccgcctc	tgccatcacc	ccgtctgtct	180
cccgcaccag	cttcacctcc	gtgtcccggg	ccgg			214

<210> 135

<211> 355

<212> DNA

<213> homo sapien

<400> 135

gaattcggca	cgaggtgaac	aggaccggtc	gccatgggccc	gtgtgatccg	tggacagagg	60
aagggcgccg	ggtctgtgtt	ccgcgcgcac	gtgaagcacc	gtaaaggcgc	tgcgcgcctg	120
cgcgccgtgg	atttcgctga	gcggcacggc	tacatcaagg	gcacgtcaa	ggacatcatc	180
cacgacccgg	gccgcggcgc	gccccctgcc	aaggtggtct	tccgggatcc	gtatcggttt	240
aagaagcgga	cggagctgtt	cattgccgcc	gagggcattc	acacgggcca	gtttgtgtat	300
tgcggcaaga	aggccagct	caacattggc	aatgtgctcc	ctgtgggcac	catgc	355

<210> 136

<211> 242

<212> DNA

<213> homo sapien

<400> 136

gaattcggca	cgagccagct	cctaaccgcg	agtgatccgc	cagcctccgc	ctcccagaggt	60
gcccgattg	cagacggagt	ctccttcact	cagtgtctca	tggtgcccag	gctggagtgc	120

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72

agtgggtgtga tctcggctcg ctacaacatc cacctcccag cagcctgcct tggcctccca 180
 aagtgccgag attgcagctc tctgcccggc cgccaccctt gtctgggaag tgaggatgct 240
 gt 242

<210> 137
 <211> 424
 <212> DNA
 <213> homo sapien

<400> 137
 gaattcggca cgagcccaga tcccagagtc cgacagcgcc cggcccagat ccccacgcct 60
 gccaggagca agccgagagc cagccggccg gcgcactccg actccgagca gtctctgtcc 120
 ttcgaccgca gccccgcgcc ctttccggga ccctgcccc gcgggcagcg ctgccaacct 180
 gccggccatg gagacccgt cccagcggcg cgccaccgc agcggggcg aggccagctc 240
 cactccgctg tcgcccacc gcatacccg gctgcaggag aaggaggacc tgcaggagct 300
 caatgatcgc ttggcggtct acatcgaccg tgtgcgctcg ctggaacgg agaacgcagg 360
 gctgcgcctt cgcataccg agtctgaaga ggtggtcagc cgcgagggtt ccggcatcaa 420
 ggcc 424

<210> 138
 <211> 448
 <212> DNA
 <213> homo sapien

<400> 138
 gaattcggca cgagcctgtg ttccaggagc cgaatcagaa atgtcatcct caggcacgcc 60
 agacttacct gtcctactca ccgatttgaa gattcaatat actaagatct tcataaacia 120
 tgaatggcat gattcagtga gtggcaagaa atttcctgtc tttaatcctg caactgagga 180
 ggagctctgc caggtagaag aaggagataa ggaggatgtt gacaaggcag tgaaggccgc 240
 aagacaggct ttccagattg gatccccgtg gcgtactatg gatgcttccg agagggggcg 300
 actattatac aagttggctg atttaatcga aagagatcgt ctgctgctgg ccgacaatgg 360
 agtcaatgaa tggaggaaaa ctctattcca atgcatatct gaatgattta gcaggctgca 420
 tcaaaacatt gcgctactgt gcagggtg 448

<210> 139
 <211> 510
 <212> DNA
 <213> homo sapien

<400> 139
 gaattcggca cgaggttccg tgcagctcac ggagaagcga atggacaaag tcggcaagta 60
 cccaaggag ctgcgcaagt gctgcgagga cggcatgcgg gagaaccca tgaggttctc 120
 gtgccagcgc cggaccggtt tcatctccct ggcgaggcgt gcaagaaggc ctctctggac 180
 tgctgcaact acatcacaga gctgcggcgg cagcacgcgc gggccagcca cctggcctgc 240
 caggagtaac ctggatgagg acatcattgc agaagagaac atcgtttccc gaagtgagtt 300
 cccagagagc tggctgtgga acgttgagga cttgaaagag ccaccgaaaa atggaatctc 360
 tacgaagctc atgaatatat ttttgaaaga ctccatcacc acgtgggaga ttctggctgt 420
 gagcatgtcg gacaagaaag ggatctgtgt ggcagacccc ttcgagggtca cagtaatgca 480
 ggacttcttc atcgacctgc ggctacccta 510

<210> 140
 <211> 360
 <212> DNA
 <213> homo sapien

<400> 140

gaattcggca	cgagcggtaa	ctaccccggc	tgcgcacagc	tggcgctcc	ttcccgtcc	60
ctcacacacc	ggcctcagcc	cgcaccggca	gtagaagatg	gtgaaagaaa	caacttacta	120
cgatgttttg	gggggtcaaac	ccaatgctac	tcaggaagaa	ttgaaaaagg	cttataggaa	180
actggctttg	aagtaccatc	ctgataagaa	cccaaataaa	ggagagaagt	ttaaacagat	240
ttctcaagct	tacgaagttc	tctctgatgc	aaagaaaagg	gaattatatg	acaaaggagg	300
agaacaggca	attaaagagg	gtggagcagg	tggcggtttt	ggctcccca	tggacatctt	360

<210> 141

<211> 483

<212> DNA

<213> homo sapien

<400> 141

gaattcggca	cgagagcaga	ggctgatctt	tgctggaaaa	cagctggaag	atgggctgca	60
ccctgtctga	ctacaacatc	cagaaagagt	ccaccctgca	cctgggtgctc	cgtctcagag	120
gtgggatgca	aatcttcgtg	aagacactca	ctggcaagac	catcaccctt	gagggtggagc	180
ccagtgcacac	catcgagaac	gtcaaagcaa	agatccagga	caaggaaggc	attcctcctg	240
accagcagag	gttgatcttt	gccggaaaagc	agctggaaga	tgggcgcacc	ctgtctgact	300
acaacatcca	gaaagagtct	accctgcacc	tggtgctccg	tctcagaggt	gggatgcaga	360
tcttcgtgaa	gaccctgact	ggtaagacca	tcaccctcga	ggtggagccc	agtgcacca	420
tcgagaatgt	caaggcaaag	atccaagata	aggaaggcat	tcctcctgat	cagcagaggt	480
tga						483

<210> 142

<211> 500

<212> DNA

<213> homo sapien

<400> 142

gaattcggca	cgagggcgcg	acgaccggcg	ggagcgtgtg	cagcggcgcg	ggcggaagtg	60
gccggcgagc	ccggtccccg	ccggcaccat	gcttcccttg	tcactgctga	agacgggtca	120
gaatcacccc	atgttggtgg	agctgaaaaa	tggggagacg	tacaatggac	acctgggtgag	180
ctgcgacaac	tggatgaaca	ttaacctgcg	agaagtcac	tgcacgtcca	gggacgggga	240
caagttctgg	cggatgcccc	agtgtcatat	ccgcggcagc	accatcaagt	acctgcgcac	300
ccccgacgag	atcatcgaca	tggtcaagga	ggaggtgggtg	gccaagggcc	gccggccggg	360
aggcctgcag	cagcagaagc	agcagaaaag	ccgcggcatg	ggcggcgctg	gccgaggtgt	420
gtttgggtggc	cggggccgag	gtgggatccc	gggcacaggc	agaagccagc	cagagaagaa	480
gcctggcaga	caggcgggca					500

<210> 143

<211> 400

<212> DNA

<213> homo sapien

<400> 143

gaattcggca	cgagctcgga	tgctcagcagg	cgtcccaacc	cagcaggaac	tggctcaatt	60
ctcagaagaa	agcgatcggc	cccgaggcag	gaaggccggc	tccgggtgcag	ggcgcgcgcg	120
ctgcgggctg	cttcggggcca	gggtcgaccc	gagggccagc	gcaagcagcg	gcaacaggag	180
cgccaggagg	acatgaggct	ctgcctgcag	tcagcaactt	ggaatattca	gacttcagac	240
cagcatcaca	gattataacc	ctccgtaaat	catctgcac	ccagctccca	tcaaaagcca	300
gcctgaagga	cccatggaca	cgtgactcca	gtgttctcaa	caacatctta	gatcaagttg	360
gtttgcacaa	catttgcac	tacttgggac	aaagcaagaa			400

<210> 144

<211> 243
 <212> DNA
 <213> homo sapien

<400> 144

gaattcggca	cgagccagct	cctaaccgcg	agtgatccgc	cagcctccgc	ctcccagaggt	60
gcccggattg	cagacggagt	ctccttcaact	cagtgtctcaa	tgggtgccag	gctggagtg	120
agtgggtgtga	tctcggctcg	ctacaacatc	cacctcccag	cagcctgcct	tggcctccca	180
aagtgccgag	attgcagcct	ctgcccggcc	gtcaccccg	ctgggaagtg	aggagcgttt	240
ctg						243

<210> 145
 <211> 450
 <212> DNA
 <213> homo sapien

<400> 145

gaattcggca	cgaggacagc	aggaccgtgg	aggccgcggc	aggggtggca	gtgggtggcgg	60
cgccggcggc	ggcgggtgtg	gttacaaccg	cagcagtggt	ggctatgaac	ccagaggtcg	120
tggaggtggc	cgtggaggca	gaggtggcat	ggcggaagt	gaccgtgggtg	gcttcaataa	180
atttgggtggc	cctcgggacc	aaggatcacg	tcatgactcc	gaacaggata	attcagacaa	240
caacaccatc	tttgtgcaag	gcctgggtga	gaatgttaca	attgagtctg	tggctgatta	300
cttcaagcag	attgggtatta	ttaagacaaa	caagaaaacg	ggacagccca	tgattaattt	360
gtacacagac	agggaaactg	gcaagctgaa	gggagaggca	acggtctctt	ttgatgacct	420
accttcagct	aaagcagcct	attgactggc				450

<210> 146
 <211> 451
 <212> DNA
 <213> homo sapien

<400> 146

gaattcggca	cgagccatcg	agtccctgcc	tttcgacttg	cagagaaatg	tctcgtgat	60
gcgggagatc	gacgcgaaat	accaagagat	cctgaaggag	ctagacgagt	gctacgagcg	120
cttcagtcgc	gagacagacg	ggcgcgagaa	gcggcggatg	ctgcactgtg	tgcagcgcg	180
gctgatccgc	accaggagct	ggcgacgag	aagatccaga	tcgtgagcca	gatgggtggag	240
ctggtggaga	accgcacgcg	gcaggtggac	agccacgtgg	agctgttcga	ggcgacgag	300
gagctgggcg	acacagcggg	caacagcggc	aaggctggcg	cggacaggcc	caaaggcgag	360
gcggcagcgc	aggctgacaa	gcccacacgc	aagcgctcac	ggcggcagcg	caacaacgag	420
aaccgtgaga	acgcgtccag	caaccacgac	c			451

<210> 147
 <211> 400
 <212> DNA
 <213> homo sapien

<400> 147

gaattcggca	cgagctcgga	tgtcagcagg	cgtcccaacc	cagcaggaac	tggctcaatt	60
ctcagaagaa	agcgatcggc	cccaggcgag	gaaggccggc	tccgggtgcag	ggcgcgccgc	120
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gcctgaagga	cccatggaca	cgtgactcca	gtgttctcaa	caacatctta	gatcaagttg	360
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<210> 148
 <211> 503
 <212> DNA
 <213> Homo sapien

<400> 148

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<400> 149

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<210> 150
 <211> 781
 <212> DNA
 <213> homo sapien

<400> 150

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<210> 151

<211> 3275

<212> DNA

<213> Homo sapien

<400> 151

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<210> 152

<211> 2179

<212> DNA

<213> homo sapien

<400> 152

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<210> 153

<211> 2109

<212> DNA

<213> Homo sapien

<400> 153

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<210> 154

<211> 1411

<212> DNA

<213> homo sapien

<400> 154

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<210> 155

<211> 678

<212> DNA

<213> homo sapien

<400> 155

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<210> 156

<211> 2668

<212> DNA

<213> Homo sapien

<400> 156

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<210> 157

<211> 2313

<212> DNA

<213> homo sapien

<400> 157

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<210> 158

<211> 2114

<212> DNA

<213> homo sapien

<400> 158

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gacacctcag	cagcgcacca	ggtggtttta	ggagaaaact	tgatagccac	agccctttgt	180
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cgggagtact	tcta					2114

<210> 159

<211> 278

<212> DNA

<213> homo sapien

<400> 159

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tgcagaatga	gaatcactcc	taaaataggt	aatggtaaaa	attaaattga	caattacctc	180
tctctatgca	gaaggaaata	tcacctatat	gacatcatca	tcctctattg	atacttgctg	240
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<210> 160

<211> 848

<212> DNA

<213> homo sapien

<400> 160

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cctagagctc	tacttacaga	acagccctga	ggcctgtgac	tatgggtctc	gaagggggca	780
ggagtcagca	ataaagctat	gtctgatatt	ttccttact	aaaaaaaaaa	aaaaaaaaaa	840
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<210> 161

<211> 432

<212> DNA

<213> homo sapien

<400> 161

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ggaagggaag	ggcccaagta	aagcacagcg	cgggagccta	gagcacatga	agctgaccc	180
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cctcctgtcc	cagcgagagc	aggaaatagt	ggcctgagc	cagcaactgc	aggaagccag	360
ggaacaagg	gagctgaagg	agcagtcact	tcagagtc	ctggatgagg	cccagagagc	420
cctagcccag	ag					432

<210> 162

<211> 433

<212> DNA

<213> homo sapien

<400> 162

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<210> 163

<211> 432

<212> DNA

<213> homo sapien

<400> 163

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gctgcaacct	gaaggggacg	cagacagtgc	cgcgggctcg	gccgtgccct	ctgagtgcct	180
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ccccgctttc	ctcacgcccc	gtccgacaaa	gcggctctcc	agcaagaagg	tggcaaggta	420
cctgcaccag	tc					432

<210> 164

<211> 395

<212> DNA

<213> homo sapien

<400> 164

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<210> 165
 <211> 503
 <212> DNA
 <213> homo sapien

<400> 165
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 agatggtgaa agaaacaact tactacgatg ttttgggggt caaacccaat gctactcagg 180
 aagaattgaa aaaggcttat aggaactgg ccttgaagta ccacctgat aagaacccaa 240
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<210> 166
 <211> 893
 <212> DNA
 <213> homo sapien

<400> 166
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 aacggaagta gttgtaggta gtggtatggt ggtatgagtc tgttttctgt tacttataac 240
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 acaataaatt ttttttggtc aaatttaaaa aaaaaaaaaa aaaaaaactc gag 893

<210> 167
 <211> 549
 <212> DNA
 <213> homo sapien

<400> 167
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 ttgacccga gccccgcgcc ctttcgggga cccctgcccc gcgggcagcg ctgccaacct 180
 gccggccatg gagaccccg cccagcggcg cggcaccgc agcggggcgc aggccagctc 240
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 gcgcgcccgc ctgcagctgg agctgagcaa agtgcgtgaa gaggtttaagg agctgaaagc 540
 gcgcaatac 549

<210> 168
 <211> 547
 <212> DNA
 <213> homo sapien

<400> 168

gaattcggca	cgagatggcg	gcaggggtcg	aagcggcggc	ggaggtggcg	gcgacggaga	60
tcaaaatgga	ggaagagagc	ggcgcgcccc	gcgtgccgag	cggcaacggg	gctccgggcc	120
ctaagggtga	aggagaacga	cctgctcaga	atgagaagag	gaaggagaaa	aacataaaaa	180
gaggaggcaa	tcgctttgag	ccatattgcca	atccaactaa	aagatacaga	gccttcatta	240
caaacatacc	ttttgatgtg	aaatggcagt	cacttaaaga	cctgggttaa	gaaaaagtgt	300
gtgaggtaac	atacgtggag	ctcttaatgg	acgctgaagg	aaagtcaagg	ggatgtgctg	360
ttgttggaatt	caagatggaa	gagagcatga	aaaaagctgc	ggaagtccta	aacaagcata	420
gtctgagcgg	aagaccactg	aaagtcaaag	aagatcctga	tggtgaacat	gccaggagag	480
caatgcaaaa	ggctggaaga	cttggaagca	cagtatttgt	agcaaatctg	gattataaag	540
ttggctg						547

<210> 169
 <211> 547
 <212> DNA
 <213> homo sapien

<400> 169

gaattcggca	ccaggagtcc	gactgtgctc	gctgctcagc	gccgcacccg	gaagatgagg	60
ctcgccgtgg	gagccctgct	ggtctgcgcc	gtcctggggc	tgtgtctggc	tgtccctgat	120
aaaactgtga	gatggtgtgc	agtgtcggag	catgaggcca	ctaagtcca	gagtttccgc	180
gaccatatga	aaagcgtcat	tccattccgat	ggtcccagtg	ttgcttgtgt	gaagaaagcc	240
tcctaccttg	attgcatcag	ggccattgcg	gcaaacgaag	cggatgctgt	gacactggat	300
gcagggtttg	tgtatgatgc	ttacctggct	cccaataacc	tgaagcctgt	ggtggcagag	360
ttctatgggt	caaaagagga	tccacagact	ttctattatg	ctgttgctgt	ggtgaagaag	420
gatagtggct	tccagatgaa	ccagcttcga	ggcaagaagt	cctgccacac	gggtctaggc	480
aggtccgctg	ggtggaacat	ccccataggc	ttactttact	gtgacttacc	tgagccacgt	540
aaacctc						547

<210> 170
 <211> 838
 <212> DNA
 <213> homo sapien

<400> 170

gaattcggca	ccagaggagc	tcggcctgcg	ctgcgccacg	atgtccgggg	agtcagccag	60
gagcttgggg	aagggaagcg	cgccccggg	gccggtcccg	gagggtctga	tccgcatcta	120
cagcatgagg	ttctgcccgt	ttgctgagag	gacgcgtcta	gtcctgaagg	ccaaggggat	180
caggcatgaa	gtcatcaata	tcaacctgaa	aaataagcct	gagtggttct	ttaagaaaaa	240
tccctttggt	ctgggtgccag	ttctggaaaa	cagtcagggt	cagctgatct	acgagtctgc	300
catcacctgt	gagtacctgg	atgaagcata	cccagggaag	aagctgttgc	cggatgaccc	360
ctatgagaaa	gcttgccaga	agatgatctt	agagtgtttt	tctaagggtg	catccttggg	420
aggaagcttt	attagaagcc	aaaataaaga	agactatgat	ggcctaaaag	aagaatttcg	480
taaagaattt	accaagctag	aggaggttct	gactaataag	aagacgacct	tctttgggtg	540
caattctatc	tctatgattg	attacctcat	ctggccctgg	tttgaacggc	tggaaagcaat	600
gaagttaaatt	gagtgtgtag	accacactcc	aaaactgaaa	ctgtggatgg	cagccatgaa	660
ggaagatccc	acagtctcag	ccctgcttac	tagtgagaaa	gactggcaag	gtttcctaga	720
gctctactta	cagaacagcc	ctgaggcctg	tgactatggg	ctctgaaggg	ggcaggagtc	780
agcaataaag	ctatgtctga	tattttcctt	cactaaaaaa	aaaaaaaaaa	aactcagag	838

<210> 171
 <211> 547
 <212> DNA
 <213> homo sapien

<400> 171
 gaattcggca ccagcgggat ttgggtcgca gttcttgttt gtggattgct gtgatcgta 60
 cttgacaatg cagatcttcg tgaagactct gactggtaag accatcacc tcgaggttga 120
 gcccagtgac accatcgaga atgtcaaggc aaagatccaa gataaggaag gcatccctcc 180
 tgaccagcag aggctgatct ttgctggaaa acagctggaa gatgggcgca ccctgtctga 240
 ctacaacatc cagaaagagt ccacctgca cctgggtgctc cgtctcagag gtgggatgca 300
 aatcttcgtg aagacactca ctggcaagac catcaccctt gaggtcgagc ccagtgacac 360
 catcgagaac gtcaaagcaa agatccagga caaggaaggc attcctctg accagcagag 420
 gttgatcttt gccggaagc agctggaaga tgggcgcacc ctgtctgact acaacatcca 480
 gaaagagtct accctgcacc tgggtgctccg tctcagaggt gggatgcaga tcttcgtgaa 540
 gaccctg 547

<210> 172
 <211> 608
 <212> DNA
 <213> homo sapien

<400> 172
 gaattcggca ccagagactt ctccctctga ggcctgcgca cccctcctca tcagcctgtc 60
 caccctcatc tacaatggtg ccctgccatg tcaagtgaac cctcaagggt cactgagttc 120
 tgagtgaac cctcatggtg gtcagtgcct gtgcaagcct ggagtgggtg ggcgcgctg 180
 tgacctctgt gcccctggct actatggctt tggccccaca ggctgtcaag gcgcttgctt 240
 gggctgcccgt gatcacacag ggggtgagca ctgtgaaagg tgcattgctg gtttccacgg 300
 ggaccacagg ctgccatag ggggccagtg ccggccctgt ccctgtcctg aaggccctgg 360
 gagccaacgg cactttgcta cttcttgcca ccaggatgaa tattcccagc agattgtgtg 420
 ccactgccgg gcaggctata cggggctgct atgtgaagct tgtgcccctg ggcactttgg 480
 ggaccatca aggccagggt gccggtgcca actgtgtgag tgcagtggga acattgaccc 540
 aatggatcct gatgcctgtg accccacac ggggcaatgc ctgcgctgtt tacaccacac 600
 agagggtc 608

<210> 173
 <211> 543
 <212> DNA
 <213> homo sapien

<400> 173
 gaattcggca ccagagatca tccgccagca gggcttgccc tctacgact acgtgcgccc 60
 ccgcctcacg gctgaggacc tgctcgaggc tcggatcatc tctctcgaga cctacaacct 120
 gctccgggag ggcaccagga gcctccgtga ggctctcgag ggcgagtcg cctgggtgcta 180
 cctctatggc acgggctccg tggctggtgt ctacctgccc ggttccaggc agacactgag 240
 catctaccag gctctcaaga aagggtgctt gagtcccgag gtggcccgcc tgctgtgga 300
 ggcacaggca gccacaggct tctgtctgga ccggtgaag ggggaacggc tgactgtgga 360
 tgaagctgtg cggaagggcc tcgtggggcc cgaactgcac gaccgctgc tctcggctga 420
 gcgggcgggt accggctacc gtgacccta caccgagcag accatctcgc tcttccaggc 480
 catgaagaag gaactgatcc ctactgagga ggcctgcgg ctgtggatgc ccagctggcc 540
 acc 543

<210> 174
 <211> 548
 <212> DNA

<213> homo sapien

<400> 174

gaattcggca cgagaaatgg cggcaggggt cgaagcggcg gcgagaggtgg cggcgacgga	60
gatcaaaatg gaggaagaga gcggcgcgcc cggcgtgccg agcggcaacg gggctccggg	120
ccctaaggggt gaaggagAAC gacctgctca gaatgagaag aggaaggaga aaacataaa	180
aagaggaggc aatcgctttg agccatatgc caatccaact aaaagataga gagccttcat	240
tacaaacata ccttttgatg tgaaatggca gtcacttaaa gacctgggta aagaaaaagt	300
tggtgaggtg acatacgtgg agctcttaat ggacgctgaa ggaaagtcaa ggggatgtgc	360
tggtgttgaa ttcaagatgg aagagagcat gaaaaaagct gcggaagtcc taaacaagca	420
tagtctgagc ggaagaccac tgaaagtcaa agaagatcct gatggtgaac atgccaggag	480
agcaatgcaa aaggtgatgg ctacgactgg tgggatgggt atgggaccag gtggcccagg	540
aatgatta	548

<210> 175

<211> 604

<212> DNA

<213> homo sapien

<400> 175

gaattcggca ccagaggacc tccaggacat gttcatcgtc cataccatcg aggagattga	60
gggcctgata tcagcccatg accagttcaa gtccaccctg ccggacgccg atagggagcg	120
cgaggccatc ctggccatcc acaaggaggc ccagaggatc gctgagagca accacatcaa	180
gctgtcgggc agcaaccctt acaccaccgt caccgcgcaa atcatcaact ccaagtggga	240
gaaggtgcag cagctggtgc caaaacggga ccatgccctc ctggaggagc agagcaagca	300
gcagtccaac gagcacctgc gccgccagtt cgccagccag gccaatgttg tggggccctg	360
gatccagacc aagatggagg agatcgggcy catctccatt gagatgaacg ggaccctgga	420
ggaccagctg agccacctga agcagtatga acgcagcatc gtggactaca agcccaacct	480
ggacctgctg gagcagcagc accagcttat ccaggaggcc ctcatcttcg acaacaagca	540
caccaactat accatggagc acatccgcgt gggctgggag cagctgctca ccaccattgc	600
ccgg	604

<210> 176

<211> 486

<212> DNA

<213> homo sapien

<400> 176

gaattcggca ccagccaagc tcaactattga atccacgccg ttcaatgtcg cagaggggaa	60
ggaggttctt ctactcgccc acaacctgcc ccagaatcgt attggttaca gctggtacaa	120
aggcgaaaga gtggatggca acagtctaatt tgtaggatat gtaataggaa ctcaacaagc	180
tacccagggt cccgcataca gtggtcgaga gacaatatac cccaatgcat ccctgctgat	240
ccagaacgtc acccagaatg acacaggatt ctatacccta caagtcataa agtcagatct	300
tgtgaatgaa gaagcaaccg gacagttcca tgtatacccg gagctgccc aagccctccat	360
ctccagcaac aactccaacc ccgtggaggga caaggatgct gtggccttca cctgtgaacc	420
tgaggttcag aacacaacct acctgtggtg ggtaaatggt cagagcctcc cggtcagtcc	480
caaggc	486

<210> 177

<211> 387

<212> DNA

<213> homo sapien

<400> 177

gaattcggca ccaggagacag cagaccagac agtcacagca gccttgacaa aacgttcctg	60
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gaactcaagc	tcttctccac	agaggaggac	agagcagaca	gcagagacca	tggagtctcc	120
ctcggccctt	ccccacagat	ggtgcatccc	ctggcagagg	ctcctgctca	cagcctcact	180
tctaaccctt	tggaaccgcg	ccaccactgc	caagctcact	attgaatcca	cgccgttcaa	240
tgtcgcagag	gggaaggagg	tgcttctact	tgccacacat	ctgccccagc	atctttttgg	300
ctacagctgg	tacaaagggtg	aaagagtggg	tggcaaccgt	caaattatag	gatatgtaat	360
aggaactcaa	caagctaccc	cagggcc				387

<210> 178

<211> 440

<212> DNA

<213> homo sapien

<400> 178

gaattcggca	cgaggagaag	cagaaaaaca	aggaatttag	ccagacttta	gaaaatgaga	60
aaaatacctt	actgagtcag	atatcaacaa	aggatgggtga	actaaaaatg	cttcaggagg	120
aagtaaccaa	aatgaacctg	ttaaatacagc	aaatccaaga	agaactctct	agagttacca	180
aactaaagga	gacagcagaa	gaagagaaaag	atgatttgga	agagaggctt	atgaatcaat	240
tagcagaact	taatggaagc	attgggaatt	actgtcagga	tggtacagat	gccccaaataa	300
aaaatgagct	attggaatct	gaaatgaaga	accttaaaaa	gtgtgtgagt	gaattggaag	360
aagaaaagca	gcagttagtc	aaggaaaaaa	ctaagggtgga	atcagaaaata	cgaaaggaat	420
atttgagaa	aatacaaggt					440

<210> 179

<211> 443

<212> DNA

<213> homo sapien

<400> 179

gaattcggca	ccagcggggg	gctacggcgg	cggtacggc	ggcgtcctga	ccgcgtccga	60
cgggctgctg	gcgggcaacg	agaagctaac	catgcagaac	ctcaacgacc	gcctggcctc	120
ctacctggac	aagggtgcgcg	ccctggaggc	ggccaacggc	gagctagagg	tgaagatccg	180
cgactggtac	cagaagcagg	ggcctgggce	eteecggagc	tacagccact	actacacgac	240
catccaggac	ctgcgggaca	agattcttgg	tgccaccatt	gagaactcca	ggattgtcct	300
gcagatcgac	aacgcccgtc	tggctgcaga	tgacttccga	accaagtttg	agacggaaca	360
ggctctgcgc	atgagcgtgg	aggccgacat	caacggcctg	cgcaggggtg	tggatgagct	420
gaccctggcc	aggaccgacc	tgg				443

<210> 180

<211> 403

<212> DNA

<213> homo sapien

<400> 180

gaattcggca	cgaggttatg	agagtcgact	tcaatgttcc	tatgaagaac	aaccagataa	60
caaacaacca	gaggattaag	gctgctgtcc	caagcatcaa	attctgcttg	gacaatggag	120
ccaagtcggt	agtccttatg	agccacctag	gccggcctga	tgggtgtccc	atgcctgaca	180
agtactcctt	agagccagtt	gctgtagaac	tcagatctct	gctgggcaag	gatgttctgt	240
tcttgaagga	ctgtgtaggc	ccagaagtgg	agaaagcctg	tgccaaccca	gctgctgggt	300
ctgtcatcct	gctggagaac	ctccgctttc	atgtggagga	agaagggaag	ggaaaagatg	360
cttctgggaa	caaggttaaa	gccgagccag	ccaaaataga	agc		403

<210> 181

<211> 493

<212> DNA

<213> homo sapien

<400> 181

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gaattcggca ccagcagagg tctccagagc cttctctctc ctgtgcaaaa tggcaactct      60
taaggaaaaa ctcattgcac cagttgcgga agaagaggca acagttccaa acaataagat      120
cactgtagtg ggtgttggaac aagttggtat ggcgtgtgct atcagcattc tgggaaagtc      180
tctggctgat gaacttgctc ttgtggatgt tttggaagat aagcttaaag gagaaatgat      240
ggatctgcag catgggagct tatttcttca gacacctaaa attgtggcag ataaagatta      300
ttctgtgacc gccaatctta agattgtagt ggtaactgca ggagtccgtc agcaagaagg      360
ggagagtcgg ctcaatctgg tgcagagaaa tgtaaatgtc ttcaaattca ttattcctca      420
gatcgtcaag tacagtcttg attgcatcat aattgtggtt tccaaccagc tggacattct      480
tacgtatgtt acc                                     493

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<210> 182

<211> 209

<212> PRT

<213> homo sapien

<400> 182

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Ala Phe Ser Ser Asn Pro Lys Val Gln Val Glu Ala Ile Glu Gly Gly
-1      5      10      15
Ala Leu Gln Lys Leu Leu Val Ile Leu Ala Thr Glu Gln Pro Leu Thr
      20      25      30
Ala Lys Lys Lys Val Leu Phe Ala Leu Cys Ser Leu Leu Arg His Phe
      35      40      45
Pro Tyr Ala Gln Arg Gln Phe Leu Lys Leu Gly Gly Leu Gln Val Leu
      50      55      60
Arg Thr Leu Val Gln Glu Lys Gly Thr Glu Val Leu Ala Val Arg Val
      65      70      75      80
Val Thr Leu Leu Tyr Asp Leu Val Thr Glu Lys Met Phe Ala Glu Glu
      85      90      95
Glu Ala Glu Leu Thr Gln Glu Met Ser Pro Glu Lys Leu Gln Gln Tyr
      100      105      110
Arg Gln Val His Leu Leu Pro Gly Leu Trp Glu Gln Gly Trp Cys Glu
      115      120      125
Ile Thr Ala His Leu Leu Ala Leu Pro Glu His Asp Ala Arg Glu Lys
      130      135      140
Val Leu Gln Thr Leu Gly Val Leu Leu Thr Thr Cys Arg Asp Arg Tyr
      145      150      155      160
Arg Gln Asp Pro Gln Leu Gly Arg Thr Leu Ala Ser Leu Gln Ala Glu
      165      170      175
Tyr Gln Val Leu Ala Ser Leu Glu Leu Gln Asp Gly Glu Asp Glu Gly
      180      185      190
Tyr Phe Gln Glu Leu Leu Gly Ser Val Asn Ser Leu Leu Lys Glu Leu
      195      200      205
Arg

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<210> 183

<211> 255

<212> PRT

<213> homo sapien

<400> 183

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Met Ala Ala Gly Val Glu Ala Ala Ala Glu Val Ala Ala Thr Glu Pro
1      5      10      15

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Lys Met Glu Glu Glu Ser Gly Ala Pro Cys Val Pro Ser Gly Asn Gly
 20 25 30
 Ala Pro Gly Pro Lys Gly Glu Glu Arg Pro Thr Gln Asn Glu Lys Arg
 35 40 45
 Lys Glu Lys Asn Ile Lys Arg Gly Gly Asn Arg Phe Glu Pro Tyr Ser
 50 55 60
 Asn Pro Thr Lys Arg Tyr Arg Ala Phe Ile Thr Asn Ile Pro Phe Asp
 65 70 75 80
 Val Lys Trp Gln Ser Leu Lys Asp Leu Val Lys Glu Lys Val Gly Glu
 85 90 95
 Val Thr Tyr Val Glu Leu Leu Met Asp Ala Glu Gly Lys Ser Arg Gly
 100 105 110
 Cys Ala Val Val Glu Phe Lys Met Glu Glu Ser Met Lys Lys Ala Ala
 115 120 125
 Glu Val Leu Asn Lys His Ser Leu Ser Gly Arg Pro Leu Lys Val Lys
 130 135 140
 Glu Asp Pro Asp Gly Glu His Ala Arg Arg Ala Met Gln Lys Ala Gly
 145 150 155 160
 Arg Leu Gly Ser Thr Val Phe Val Ala Asn Leu Asp Tyr Lys Val Gly
 165 170 175
 Trp Lys Lys Leu Lys Glu Val Phe Ser Met Ala Gly Val Val Val Arg
 180 185 190
 Ala Asp Ile Leu Glu Asp Lys Asp Gly Lys Ser Arg Gly Ile Gly Ile
 195 200 205
 Val Thr Phe Glu Gln Ser Ile Glu Ala Val Gln Ala Ile Ser Met Phe
 210 215 220
 Asn Gly Gln Leu Leu Phe Asp Arg Pro Met His Val Lys Met Asp Glu
 225 230 235 240
 Arg Ala Leu Pro Lys Gly Asp Phe Phe Pro Pro Glu Arg His Ser
 245 250 255

<210> 184

<211> 188

<212> PRT

<213> Homo sapien

<400> 184

Leu Ser Gly Ser Cys Ile Arg Arg Glu Gln Thr Pro Glu Lys Glu Lys
 1 5 10 15
 Gln Val Val Leu Phe Glu Glu Ala Ser Trp Thr Cys Thr Pro Ala Cys
 20 25 30
 Gly Asp Glu Pro Arg Thr Val Ile Leu Leu Ser Ser Met Leu Ala Asp
 35 40 45
 His Arg Leu Lys Leu Glu Asp Tyr Lys Asp Arg Leu Lys Ser Gly Glu
 50 55 60
 His Leu Asn Pro Asp Gln Leu Glu Ala Val Glu Lys Tyr Glu Glu Val
 65 70 75 80
 Leu His Asn Leu Glu Phe Ala Lys Glu Leu Gln Lys Thr Phe Ser Gly
 85 90 95
 Leu Ser Leu Asp Leu Leu Lys Ala Gln Lys Lys Ala Gln Arg Arg Glu
 100 105 110
 His Met Leu Lys Leu Glu Ala Glu Lys Lys Lys Leu Arg Thr Ile Leu
 115 120 125
 Gln Val Gln Tyr Val Leu Gln Asn Leu Thr Gln Glu His Val Gln Lys
 130 135 140

Asp Phe Lys Gly Gly Leu Asn Gly Ala Val Tyr Leu Pro Ser Lys Glu
 145 150 155 160
 Leu Asp Tyr Leu Ile Lys Phe Ser Lys Leu Thr Cys Pro Glu Arg Asn
 165 170 175
 Glu Ser Leu Arg Gln Thr Leu Glu Gly Ser Thr Val
 180 185

<210> 185

<211> 746

<212> PRT

<213> Homo sapien

<400> 185

Asp Lys His Leu Lys Asp Leu Leu Ser Lys Leu Leu Asn Ser Gly Tyr
 1 5 10 15
 Phe Glu Ser Ile Pro Val Pro Lys Asn Ala Lys Glu Lys Glu Val Pro
 20 25 30
 Leu Glu Glu Met Leu Ile Gln Ser Glu Lys Lys Thr Gln Leu Ser
 35 40 45
 Lys Thr Glu Ser Val Lys Glu Ser Glu Ser Leu Met Glu Phe Ala Gln
 50 55 60
 Pro Glu Ile Gln Pro Gln Glu Phe Leu Asn Arg Arg Tyr Met Thr Glu
 65 70 75 80
 Val Asp Tyr Ser Asn Lys Gln Gly Glu Glu Gln Pro Trp Glu Ala Asp
 85 90 95
 Tyr Ala Arg Lys Pro Asn Leu Pro Lys Arg Trp Asp Met Leu Thr Glu
 100 105 110
 Pro Asp Gly Gln Glu Lys Lys Gln Glu Ser Phe Lys Ser Trp Glu Ala
 115 120 125
 Ser Gly Lys His Gln Glu Val Ser Lys Pro Ala Val Ser Leu Glu Gln
 130 135 140
 Arg Lys Gln Asp Thr Ser Lys Leu Arg Ser Thr Leu Pro Glu Glu Gln
 145 150 155 160
 Lys Lys Gln Glu Ile Ser Lys Ser Lys Pro Ser Pro Ser Gln Trp Lys
 165 170 175
 Gln Asp Thr Pro Lys Ser Lys Ala Gly Tyr Val Gln Glu Glu Lys
 180 185 190
 Lys Gln Glu Thr Pro Lys Leu Trp Pro Val Gln Leu Gln Lys Glu Gln
 195 200 205
 Asp Pro Lys Lys Gln Thr Pro Lys Ser Trp Thr Pro Ser Met Gln Ser
 210 215 220
 Glu Gln Asn Thr Thr Lys Ser Trp Thr Thr Pro Met Cys Glu Glu Gln
 225 230 235 240
 Asp Ser Lys Gln Pro Glu Thr Pro Lys Ser Trp Glu Asn Asn Val Glu
 245 250 255
 Ser Gln Lys His Ser Leu Thr Ser Gln Ser Gln Ile Ser Pro Lys Ser
 260 265 270
 Trp Gly Val Ala Thr Ala Ser Leu Ile Pro Asn Asp Gln Leu Leu Pro
 275 280 285
 Arg Lys Leu Asn Thr Glu Pro Lys Asp Val Pro Lys Pro Val His Gln
 290 295 300
 Pro Val Gly Ser Ser Ser Thr Leu Pro Lys Asp Pro Val Leu Arg Lys
 305 310 315 320
 Glu Lys Leu Gln Asp Leu Met Thr Gln Ile Gln Gly Thr Cys Asn Phe
 325 330 335

Met Gln Glu Ser Val Leu Asp Phe Asp Lys Pro Ser Ser Ala Ile Pro
 340 345 350
 Thr Ser Gln Pro Pro Ser Ala Thr Pro Gly Ser Pro Val Ala Ser Lys
 355 360 365
 Glu Gln Asn Leu Ser Ser Gln Ser Asp Phe Leu Gln Glu Pro Leu Gln
 370 375 380
 Val Phe Asn Val Asn Ala Pro Leu Pro Pro Arg Lys Glu Gln Glu Ile
 385 390 395 400
 Lys Glu Ser Pro Tyr Ser Pro Gly Tyr Asn Gln Ser Phe Thr Thr Ala
 405 410 415
 Ser Thr Gln Thr Pro Pro Gln Cys Gln Leu Pro Ser Ile His Val Glu
 420 425 430
 Gln Thr Val His Ser Gln Glu Thr Ala Ala Asn Tyr His Pro Asp Gly
 435 440 445
 Thr Ile Gln Val Ser Asn Gly Ser Leu Ala Phe Tyr Pro Ala Gln Thr
 450 455 460
 Asn Val Phe Pro Arg Pro Thr Gln Pro Phe Val Asn Ser Arg Gly Ser
 465 470 475 480
 Val Arg Gly Cys Thr Arg Gly Gly Arg Leu Ile Thr Asn Ser Tyr Arg
 485 490 495
 Ser Pro Gly Gly Tyr Lys Gly Phe Asp Thr Tyr Arg Gly Leu Pro Ser
 500 505 510
 Ile Ser Asn Gly Asn Tyr Ser Gln Leu Gln Phe Gln Ala Arg Glu Tyr
 515 520 525
 Ser Gly Ala Pro Tyr Ser Gln Arg Asp Asn Phe Gln Gln Cys Tyr Lys
 530 535 540
 Arg Gly Gly Thr Ser Gly Gly Pro Arg Ala Asn Ser Arg Ala Gly Trp
 545 550 555 560
 Ser Asp Ser Ser Gln Val Ser Ser Pro Glu Arg Asp Asn Glu Thr Phe
 565 570 575
 Asn Ser Gly Asp Ser Gly Gln Gly Asp Ser Arg Ser Met Thr Pro Val
 580 585 590
 Asp Val Pro Val Thr Asn Pro Ala Ala Thr Ile Leu Pro Val His Val
 595 600 605
 Tyr Pro Leu Pro Gln Gln Met Arg Val Ala Phe Ser Ala Ala Arg Thr
 610 615 620
 Ser Asn Leu Ala Pro Gly Thr Leu Asp Gln Pro Ile Val Phe Asp Leu
 625 630 635 640
 Leu Leu Asn Asn Leu Gly Glu Thr Phe Asp Leu Gln Leu Gly Arg Phe
 645 650 655
 Asn Cys Pro Val Asn Gly Thr Tyr Val Phe Ile Phe His Met Leu Lys
 660 665 670
 Leu Ala Val Asn Val Pro Leu Tyr Val Asn Leu Met Lys Asn Glu Glu
 675 680 685
 Val Leu Val Ser Ala Tyr Ala Asn Asp Gly Ala Pro Asp His Glu Thr
 690 695 700
 Ala Ser Asn His Ala Ile Leu Gln Leu Phe Gln Gly Asp Gln Ile Trp
 705 710 715 720
 Leu Arg Leu His Arg Gly Ala Ile Tyr Gly Ser Ser Trp Lys Tyr Ser
 725 730 735
 Thr Phe Ser Gly Tyr Leu Leu Tyr Gln Asp
 740 745

<210> 186

<211> 705

<212> PRT

<213> Homo sapien

<400> 186

Ala Leu Leu Asn Val Arg Gln Pro Pro Ser Thr Thr Thr Phe Val Leu
 1 5 10 15
 Asn Gln Ile Asn His Leu Pro Pro Leu Gly Ser Thr Ile Val Met Thr
 20 25 30
 Lys Thr Pro Pro Val Thr Thr Asn Arg Gln Thr Ile Thr Leu Thr Lys
 35 40 45
 Phe Ile Gln Thr Thr Ala Ser Thr Arg Pro Ser Val Ser Ala Pro Thr
 50 55 60
 Val Arg Asn Ala Met Thr Ser Ala Pro Ser Lys Asp Gln Val Gln Leu
 65 70 75 80
 Lys Asp Leu Leu Lys Asn Asn Ser Leu Asn Glu Leu Met Lys Leu Lys
 85 90 95
 Pro Pro Ala Asn Ile Ala Gln Pro Val Ala Thr Ala Ala Thr Asp Val
 100 105 110
 Ser Asn Gly Thr Val Lys Lys Glu Ser Ser Asn Lys Glu Gly Ala Arg
 115 120 125
 Met Trp Ile Asn Asp Met Lys Met Arg Ser Phe Ser Pro Thr Met Lys
 130 135 140
 Val Pro Val Val Lys Glu Asp Asp Glu Pro Glu Glu Glu Asp Glu Glu
 145 150 155 160
 Glu Met Gly His Ala Glu Thr Tyr Ala Glu Tyr Met Pro Ile Lys Leu
 165 170 175
 Lys Ile Gly Leu Arg His Pro Asp Ala Val Val Glu Thr Ser Ser Leu
 180 185 190
 Ser Ser Val Thr Pro Pro Asp Val Trp Tyr Lys Thr Ser Ile Ser Glu
 195 200 205
 Glu Thr Ile Asp Asn Gly Trp Leu Ser Ala Leu Gln Leu Glu Ala Ile
 210 215 220
 Thr Tyr Ala Ala Gln Gln His Glu Thr Phe Leu Pro Asn Gly Asp Arg
 225 230 235 240
 Ala Gly Phe Leu Ile Gly Asp Gly Ala Gly Val Gly Lys Gly Arg Thr
 245 250 255
 Ile Ala Gly Ile Ile Tyr Glu Asn Tyr Leu Leu Ser Arg Lys Arg Ala
 260 265 270
 Leu Trp Phe Ser Val Ser Asn Asp Leu Lys Tyr Asp Ala Glu Arg Asp
 275 280 285
 Leu Arg Asp Ile Gly Ala Lys Asn Ile Leu Val His Ser Leu Asn Lys
 290 295 300
 Phe Lys Tyr Gly Lys Ile Ser Ser Lys His Asn Gly Ser Val Lys Lys
 305 310 315 320
 Gly Val Ile Phe Ala Thr Tyr Ser Ser Leu Ile Gly Glu Ser Gln Ser
 325 330 335
 Gly Gly Lys Tyr Lys Thr Arg Leu Lys Gln Leu Leu His Trp Cys Gly
 340 345 350
 Asp Asp Phe Asp Gly Val Ile Val Phe Asp Glu Cys His Lys Ala Lys
 355 360 365
 Asn Leu Cys Pro Val Gly Ser Ser Lys Pro Thr Lys Thr Gly Leu Ala
 370 375 380
 Val Leu Glu Leu Gln Asn Lys Leu Pro Lys Ala Arg Val Val Tyr Ala
 385 390 395 400
 Ser Ala Thr Gly Ala Ser Glu Pro Arg Asn Met Ala Tyr Met Asn Arg

405 410 415
 Leu Gly Ile Trp Gly Glu Gly Thr Pro Phe Arg Glu Phe Ser Asp Phe
 420 425 430
 Ile Gln Ala Val Glu Arg Arg Gly Val Gly Ala Met Glu Ile Val Ala
 435 440 445
 Met Asp Met Lys Leu Arg Gly Met Tyr Ile Ala Arg Gln Leu Ser Phe
 450 455 460
 Thr Gly Val Thr Phe Lys Ile Glu Glu Val Leu Leu Ser Gln Ser Tyr
 465 470 475 480
 Val Lys Met Tyr Asn Lys Ala Val Lys Leu Trp Val Ile Ala Arg Glu
 485 490 495
 Arg Phe Gln Gln Ala Ala Asp Leu Ile Asp Ala Glu Gln Arg Met Lys
 500 505 510
 Lys Ser Met Trp Gly Gln Phe Trp Ser Ala His Gln Arg Phe Phe Lys
 515 520 525
 Tyr Leu Cys Ile Ala Ser Lys Val Lys Arg Val Val Gln Leu Ala Arg
 530 535 540
 Glu Glu Ile Lys Asn Gly Lys Cys Val Val Ile Gly Leu Gln Ser Thr
 545 550 555 560
 Gly Glu Ala Arg Thr Leu Glu Ala Leu Glu Gly Gly Gly Glu Leu
 565 570 575
 Asn Asp Phe Val Ser Thr Ala Lys Gly Val Leu Gln Ser Leu Ile Glu
 580 585 590
 Lys His Phe Pro Ala Pro Asp Arg Lys Lys Leu Tyr Ser Leu Leu Gly
 595 600 605
 Ile Asp Leu Thr Ala Pro Ser Asn Asn Ser Ser Pro Arg Asp Ser Pro
 610 615 620
 Cys Lys Glu Asn Lys Ile Lys Lys Arg Lys Gly Glu Glu Ile Thr Arg
 625 630 635 640
 Glu Ala Lys Lys Ala Arg Lys Val Gly Gly Leu Thr Gly Ser Ser Ser
 645 650 655
 Asp Asp Ser Gly Ser Glu Ser Asp Ala Ser Asp Asn Glu Glu Ser Asp
 660 665 670
 Tyr Glu Ser Ser Lys Asn Met Ser Ser Gly Asp Asp Asp Phe Asn
 675 680 685
 Pro Phe Leu Asp Glu Ser Asn Glu Asp Asp Glu Asn Asp Pro Trp Leu
 690 695 700
 Ile
 705

<210> 187

<211> 595

<212> PRT

<213> Homo sapien

<400> 187

Glu Ser Pro Arg His Arg Gly Glu Gly Gly Gly Glu Trp Gly Pro Gly
 1 5 10 15
 Val Pro Arg Glu Arg Arg Glu Ser Ala Gly Glu Trp Gly Ala Asp Thr
 20 25 30
 Pro Lys Glu Gly Gly Glu Ser Ala Gly Glu Trp Gly Ala Glu Val Pro
 35 40 45
 Arg Gly Arg Gly Glu Gly Ala Gly Glu Trp Gly Pro Asp Thr Pro Lys
 50 55 60
 Glu Arg Gly Gln Gly Val Arg Glu Trp Gly Pro Glu Ile Pro Gln Glu

Arg Pro Gly Pro Arg Arg Pro Ala Arg Arg Pro Arg Gly Glu Leu Ile
 515 520 525
 Pro Arg Arg Pro Asp Pro Ala Ala Pro Ser Glu Glu Gly Leu Arg Met
 530 535 540
 Glu Ser Ser Val Asp Asp Gly Ala Thr Ala Thr Thr Ala Asp Ala Ala
 545 550 555 560
 Ser Gly Glu Ala Pro Glu Ala Gly Pro Ser Pro Ser His Ser Pro Thr
 565 570 575
 Met Cys Gln Thr Gly Gly Pro Gly Pro Pro Pro Gln Pro Pro Arg
 580 585 590
 Trp Leu Pro
 595

<210> 188
 <211> 376
 <212> PRT
 <213> Homo sapien

<400> 188
 Glu Met Arg Lys Phe Asp Val Pro Ser Met Glu Ser Thr Leu Asn Gln
 1 5 10 15
 Pro Ala Met Leu Glu Thr Leu Tyr Ser Asp Pro His Tyr Arg Ala His
 20 25 30
 Phe Pro Asn Pro Arg Pro Asp Thr Asn Lys Asp Val Tyr Lys Val Leu
 35 40 45
 Pro Glu Ser Lys Lys Ala Pro Gly Ser Gly Ala Val Phe Glu Arg Asn
 50 55 60
 Gly Pro His Ala Ser Ser Ser Gly Val Leu Pro Leu Gly Leu Gln Pro
 65 70 75 80
 Ala Pro Gly Leu Ser Lys Ser Leu Ser Ser Gln Val Trp Gln Pro Ser
 85 90 95
 Pro Asp Pro Trp His Pro Gly Glu Gln Ser Cys Glu Leu Ser Thr Cys
 100 105 110
 Arg Gln Gln Leu Glu Leu Ile Arg Leu Gln Met Glu Gln Met Gln Leu
 115 120 125
 Gln Asn Gly Ala Met Cys His His Pro Ala Ala Phe Ala Pro Leu Leu
 130 135 140
 Pro Thr Leu Glu Pro Ala Gln Trp Leu Ser Ile Leu Asn Ser Asn Glu
 145 150 155 160
 His Leu Leu Lys Glu Lys Glu Leu Leu Ile Asp Lys Gln Arg Lys His
 165 170 175
 Ile Ser Gln Leu Glu Gln Lys Val Arg Glu Ser Glu Leu Gln Val His
 180 185 190
 Ser Ala Leu Leu Gly Arg Pro Ala Pro Phe Gly Asp Val Cys Leu Leu
 195 200 205
 Arg Leu Gln Glu Leu Gln Arg Glu Asn Thr Phe Leu Arg Ala Gln Phe
 210 215 220
 Ala Gln Lys Thr Glu Ala Leu Ser Lys Glu Lys Met Glu Leu Glu Lys
 225 230 235 240
 Lys Leu Ser Ala Ser Glu Val Glu Ile Gln Leu Ile Arg Glu Ser Leu
 245 250 255
 Lys Val Thr Leu Gln Lys His Ser Glu Glu Gly Lys Lys Gln Glu Glu
 260 265 270
 Arg Val Lys Gly Arg Asp Lys His Ile Asn Asn Leu Lys Lys Cys
 275 280 285

Gln Lys Glu Ser Glu Gln Asn Arg Glu Lys Gln Gln Arg Ile Glu Thr
 290 295 300
 Leu Glu Arg Tyr Leu Ala Asp Leu Pro Thr Leu Glu Asp His Gln Lys
 305 310 315 320
 Gln Thr Glu Gln Leu Lys Asp Ala Glu Leu Lys Asn Thr Glu Leu Gln
 325 330 335
 Glu Arg Val Ala Glu Leu Glu Thr Leu Leu Glu Asp Thr Gln Ala Thr
 340 345 350
 Cys Arg Glu Lys Glu Val Gln Leu Glu Ser Leu Arg Gln Arg Glu Ala
 355 360 365
 Asp Leu Ser Ser Ala Arg His Arg
 370 375

<210> 189
 <211> 160
 <212> PRT
 <213> Homo sapien

<400> 189
 Met Leu Glu Ala His Arg Arg Gln Arg His Pro Phe Leu Leu Leu Gly
 1 5 10 15
 Thr Thr Ala Asn Arg Thr Gln Ser Leu Asn Tyr Gly Cys Ile Val Glu
 20 25 30
 Asn Pro Gln Thr His Glu Val Leu His Tyr Val Glu Lys Pro Ser Thr
 35 40 45
 Phe Ile Ser Asp Ile Ile Asn Cys Gly Ile Tyr Leu Phe Ser Pro Glu
 50 55 60
 Ala Leu Lys Pro Leu Arg Asp Val Phe Gln Arg Asn Gln Gln Asp Gly
 65 70 75 80
 Gln Leu Glu Asp Ser Pro Gly Leu Trp Pro Gly Ala Gly Thr Ile Arg
 85 90 95
 Leu Glu Gln Asp Val Phe Ser Ala Leu Ala Gly Gln Gly Gln Ile Tyr
 100 105 110
 Val His Leu Thr Asp Gly Ile Trp Ser Gln Ile Lys Ser Ala Gly Ser
 115 120 125
 Ala Leu Tyr Ala Ser Arg Leu Tyr Leu Ser Arg Tyr Gln Asp Thr His
 130 135 140
 Pro Glu Arg Leu Ala Lys His Thr Pro Gly Gly Pro Trp Ile Arg Gly
 145 150 155 160

<210> 190
 <211> 146
 <212> PRT
 <213> Homo sapien

<400> 190
 Met Asp Pro Arg Ala Ser Leu Leu Leu Leu Gly Asn Val Tyr Ile His
 1 5 10 15
 Pro Thr Ala Lys Val Ala Pro Ser Ala Val Leu Gly Pro Asn Val Ser
 20 25 30
 Ile Gly Lys Gly Val Thr Val Gly Glu Gly Val Arg Leu Arg Glu Ser
 35 40 45
 Ile Val Leu His Gly Ala Thr Leu Gln Glu His Thr Cys Val Leu His
 50 55 60
 Ser Ile Val Gly Trp Gly Ser Thr Val Gly Arg Trp Ala Arg Val Glu

<210> 191
<211> 704
<212> PRT
<213> Homo sapien

<400> 191

Glu	Gly	Gly	Cys	Ala	Ala	Gly	Arg	Gly	Arg	Glu	Leu	Glu	Pro	Glu	Leu
1				5					10					15	
Glu	Pro	Gly	Pro	Gly	Pro	Gly	Ser	Ala	Leu	Glu	Pro	Gly	Glu	Glu	Phe
		20						25					30		
Glu	Ile	Val	Asp	Arg	Ser	Gln	Leu	Pro	Gly	Pro	Gly	Asp	Leu	Arg	Ser
	35					40						45			
Ala	Thr	Arg	Pro	Arg	Ala	Ala	Glu	Gly	Trp	Ser	Ala	Pro	Ile	Leu	Thr
	50					55					60				
Leu	Ala	Arg	Arg	Ala	Thr	Gly	Asn	Leu	Ser	Ala	Ser	Cys	Gly	Ser	Ala
65				70						75				80	
Leu	Arg	Ala	Ala	Ala	Gly	Leu	Gly	Gly	Gly	Asp	Ser	Gly	Asp	Gly	Thr
			85					90						95	
Ala	Arg	Ala	Ala	Ser	Lys	Cys	Gln	Met	Met	Glu	Glu	Arg	Ala	Asn	Leu
		100						105					110		
Met	His	Met	Met	Lys	Leu	Ser	Ile	Lys	Val	Leu	Leu	Gln	Ser	Ala	Leu
	115					120						125			
Ser	Leu	Gly	Arg	Ser	Leu	Asp	Ala	Asp	His	Ala	Pro	Leu	Gln	Gln	Phe
	130					135					140				
Phe	Val	Val	Met	Glu	His	Cys	Leu	Lys	His	Gly	Leu	Lys	Val	Lys	Lys
145				150						155				160	
Ser	Phe	Ile	Gly	Gln	Asn	Lys	Ser	Phe	Phe	Gly	Pro	Leu	Glu	Leu	Val
			165						170					175	
Glu	Lys	Leu	Cys	Pro	Glu	Ala	Ser	Asp	Ile	Ala	Thr	Ser	Val	Arg	Asn
		180						185					190		
Leu	Pro	Glu	Leu	Lys	Thr	Ala	Val	Gly	Arg	Gly	Arg	Ala	Trp	Leu	Tyr
	195					200						205			
Leu	Ala	Leu	Met	Gln	Lys	Lys	Leu	Ala	Asp	Tyr	Leu	Lys	Val	Leu	Ile
	210					215					220				
Asp	Asn	Lys	His	Leu	Leu	Ser	Glu	Phe	Tyr	Glu	Pro	Glu	Ala	Leu	Met
225				230						235				240	
Met	Glu	Glu	Glu	Gly	Met	Val	Ile	Val	Gly	Leu	Leu	Val	Gly	Leu	Asn
				245					250					255	
Val	Leu	Asp	Ala	Asn	Leu	Cys	Leu	Lys	Gly	Glu	Asp	Leu	Asp	Ser	Gln
		260						265					270		
Val	Gly	Val	Ile	Asp	Phe	Ser	Leu	Tyr	Leu	Lys	Asp	Val	Gln	Asp	Leu
	275					280						285			
Asp	Gly	Gly	Lys	Glu	His	Glu	Arg	Ile	Thr	Asp	Val	Leu	Asp	Gln	Lys

290 Asn Tyr Val Glu Glu Leu Asn Arg His Leu Ser Cys Thr Val Gly Asp
 305 310 315 320
 Leu Gln Thr Lys Ile Asp Gly Leu Glu Lys Thr Asn Ser Lys Leu Gln
 325 330 335
 Glu Glu Leu Ser Ala Ala Thr Asp Arg Ile Cys Ser Leu Gln Glu Glu
 340 345 350
 Gln Gln Gln Leu Arg Glu Gln Asn Glu Leu Ile Arg Glu Arg Ser Glu
 355 360 365
 Lys Ser Val Glu Ile Thr Lys Gln Asp Thr Lys Val Glu Leu Glu Thr
 370 375 380
 Tyr Lys Gln Thr Arg Gln Gly Leu Asp Glu Met Tyr Ser Asp Val Trp
 385 390 395 400
 Lys Gln Leu Lys Glu Glu Lys Lys Val Arg Leu Glu Leu Glu Lys Glu
 405 410 415
 Leu Glu Leu Gln Ile Gly Met Lys Thr Glu Met Glu Ile Ala Met Lys
 420 425 430
 Leu Leu Glu Lys Asp Thr His Glu Lys Gln Asp Thr Leu Val Ala Leu
 435 440 445
 Arg Gln Gln Leu Glu Glu Val Lys Ala Ile Asn Leu Gln Met Phe His
 450 455 460
 Lys Ala Gln Asn Ala Glu Ser Ser Leu Gln Gln Lys Asn Glu Ala Ile
 465 470 475 480
 Thr Ser Phe Glu Gly Lys Thr Asn Gln Val Met Ser Ser Met Lys Gln
 485 490 495
 Met Glu Glu Arg Leu Gln His Ser Glu Arg Ala Arg Gln Gly Ala Glu
 500 505 510
 Glu Arg Ser His Lys Leu Gln Gln Glu Leu Gly Gly Arg Ile Gly Ala
 515 520 525
 Leu Gln Leu Gln Leu Ser Gln Leu His Glu Gln Cys Ser Ser Leu Glu
 530 535 540
 Lys Glu Leu Lys Ser Glu Lys Glu Gln Arg Gln Ala Leu Gln Arg Glu
 545 550 555 560
 Leu Gln His Glu Lys Asp Thr Ser Ser Leu Leu Arg Met Glu Leu Gln
 565 570 575
 Gln Val Glu Gly Leu Lys Lys Glu Leu Arg Glu Leu Gln Asp Glu Lys
 580 585 590
 Ala Glu Leu Gln Lys Ile Cys Glu Glu Gln Glu Gln Ala Leu Gln Glu
 595 600 605
 Met Gly Leu His Leu Ser Gln Ser Lys Leu Lys Met Glu Asp Ile Lys
 610 615 620
 Glu Val Asn Gln Ala Leu Lys Gly His Ala Trp Leu Lys Asp Asp Glu
 625 630 635 640
 Ala Thr His Cys Arg Gln Cys Glu Lys Glu Phe Ser Ile Ser Arg Arg
 645 650 655
 Lys His His Cys Arg Asn Cys Gly His Ile Phe Cys Asn Thr Cys Ser
 660 665 670
 Ser Asn Glu Leu Ala Leu Pro Ser Tyr Pro Lys Pro Val Arg Val Cys
 675 680 685
 Asp Ser Cys His Thr Leu Leu Leu Gln Arg Cys Ser Ser Thr Ala Ser
 690 695 700

<210> 192

<211> 331

<212> PRT

100

<213> Homo sapien

<400> 192

Arg Ala Gly Ala Ser Ala Met Ala Leu Arg Lys Glu Leu Leu Lys Ser
 1 5 10 15
 Ile Trp Tyr Ala Phe Thr Ala Leu Asp Val Glu Lys Ser Gly Lys Val
 20 25 30
 Ser Lys Ser Gln Leu Lys Val Leu Ser His Asn Leu Tyr Thr Val Leu
 35 40 45
 His Ile Pro His Asp Pro Val Ala Leu Glu Glu His Phe Arg Asp Asp
 50 55 60
 Asp Asp Gly Pro Val Ser Ser Gln Gly Tyr Met Pro Tyr Leu Asn Lys
 65 70 75 80
 Tyr Ile Leu Asp Lys Val Glu Glu Gly Ala Phe Val Lys Glu His Phe
 85 90 95
 Asp Glu Leu Cys Trp Thr Leu Thr Ala Lys Lys Asn Tyr Arg Ala Asp
 100 105 110
 Ser Asn Gly Asn Ser Met Leu Ser Asn Gln Asp Ala Phe Arg Leu Trp
 115 120 125
 Cys Leu Phe Asn Phe Leu Ser Glu Asp Lys Tyr Pro Leu Ile Met Val
 130 135 140
 Pro Asp Glu Val Glu Tyr Leu Leu Lys Lys Val Leu Ser Ser Met Ser
 145 150 155 160
 Leu Glu Val Ser Leu Gly Glu Leu Glu Glu Leu Leu Ala Gln Glu Ala
 165 170 175
 Gln Val Ala Gln Thr Thr Gly Gly Leu Ser Val Trp Gln Phe Leu Glu
 180 185 190
 Leu Phe Asn Ser Gly Arg Cys Leu Arg Gly Val Gly Arg Asp Thr Leu
 195 200 205
 Ser Met Ala Ile His Glu Val Tyr Gln Glu Leu Ile Gln Asp Val Leu
 210 215 220
 Lys Gln Gly Tyr Leu Trp Lys Arg Gly His Leu Arg Arg Asn Trp Ala
 225 230 235 240
 Glu Arg Trp Phe Gln Leu Gln Pro Ser Cys Leu Cys Tyr Phe Gly Ser
 245 250 255
 Glu Glu Cys Lys Glu Lys Arg Gly Ile Ile Pro Leu Asp Ala His Cys
 260 265 270
 Cys Val Glu Val Leu Pro Asp Arg Asp Gly Lys Arg Cys Met Phe Cys
 275 280 285
 Val Lys Thr Ala Thr Arg Thr Tyr Glu Met Ser Ala Ser Asp Thr Arg
 290 295 300
 Gln Arg Gln Glu Trp Thr Ala Ala Ile Gln Met Ala Ile Arg Leu Gln
 305 310 315 320
 Ala Glu Gly Lys Thr Ser Leu His Lys Asp Leu
 325 330

<210> 193

<211> 475

<212> PRT

<213> Homo sapien

<400> 193

Lys Asn Ser Pro Leu Leu Ser Val Ser Ser Gln Thr Ile Thr Lys Glu
 1 5 10 15
 Asn Asn Arg Asn Val His Leu Glu His Ser Glu Gln Asn Pro Gly Ser

Arg Asn Ser Pro Gly Leu Gly Ser Leu Val Ser
 465 470 475

<210> 194
 <211> 241
 <212> PRT
 <213> Homo sapien

<400> 194
 Met Ser Gly Glu Ser Ala Arg Ser Leu Gly Lys Gly Ser Ala Pro Pro
 1 5 10 15
 Gly Pro Val Pro Glu Gly Ser Ile Arg Ile Tyr Ser Met Arg Phe Cys
 20 25 30
 Pro Phe Ala Glu Arg Thr Arg Leu Val Leu Lys Ala Lys Gly Ile Arg
 35 40 45
 His Glu Val Ile Asn Ile Asn Leu Lys Asn Lys Pro Glu Trp Phe Phe
 50 55 60
 Lys Lys Asn Pro Phe Gly Leu Val Pro Val Leu Glu Asn Ser Gln Gly
 65 70 75 80
 Gln Leu Ile Tyr Glu Ser Ala Ile Thr Cys Glu Tyr Leu Asp Glu Ala
 85 90 95
 Tyr Pro Gly Lys Lys Leu Leu Pro Asp Asp Pro Tyr Glu Lys Ala Cys
 100 105 110
 Gln Lys Met Ile Leu Glu Leu Phe Ser Lys Val Pro Ser Leu Val Gly
 115 120 125
 Ser Phe Ile Arg Ser Gln Asn Lys Glu Asp Tyr Ala Gly Leu Lys Glu
 130 135 140
 Glu Phe Arg Lys Glu Phe Thr Lys Leu Glu Glu Val Leu Thr Asn Lys
 145 150 155 160
 Lys Thr Thr Phe Phe Gly Gly Asn Ser Ile Ser Met Ile Asp Tyr Leu
 165 170 175
 Ile Trp Pro Trp Phe Glu Arg Leu Glu Ala Met Lys Leu Asn Glu Cys
 180 185 190
 Val Asp His Thr Pro Lys Leu Lys Leu Trp Met Ala Ala Met Lys Glu
 195 200 205
 Asp Pro Thr Val Ser Ala Leu Leu Thr Ser Glu Lys Asp Trp Gln Gly
 210 215 220
 Phe Leu Glu Leu Tyr Leu Gln Asn Ser Pro Glu Ala Cys Asp Tyr Gly
 225 230 235 240
 Leu

<210> 195
 <211> 138
 <212> PRT
 <213> Homo sapien

<400> 195
 Gln Thr Lys Ile Leu Glu Glu Asp Leu Glu Gln Ile Lys Leu Ser Leu
 1 5 10 15
 Arg Glu Arg Gly Arg Glu Leu Thr Thr Gln Arg Gln Leu Met Gln Glu
 20 25 30
 Arg Ala Glu Glu Gly Lys Gly Pro Ser Lys Ala Gln Arg Gly Ser Leu
 35 40 45
 Glu His Met Lys Leu Ile Leu Arg Asp Lys Glu Lys Glu Val Glu Cys

50 55 60
 Gln Gln Glu His Ile His Glu Leu Gln Glu Leu Lys Asp Gln Leu Glu
 65 70 75 80
 Gln Gln Leu Gln Gly Leu His Arg Lys Val Gly Glu Thr Ser Leu Leu
 85 90 95
 Leu Ser Gln Arg Glu Gln Glu Ile Val Leu Gln Gln Gln Leu Gln
 100 105 110
 Glu Ala Arg Glu Gln Gly Glu Leu Lys Glu Gln Ser Leu Gln Ser Gln
 115 120 125
 Leu Asp Glu Ala Gln Arg Ala Leu Ala Gln
 130 135

<210> 196

<211> 102

<212> PRT

<213> Homo sapien

<400> 196

Met Ser Lys Arg Lys Ala Pro Gln Glu Thr Leu Asn Gly Gly Ile Thr
 1 5 10 15
 Asp Met Leu Thr Glu Leu Ala Asn Phe Glu Lys Asn Val Ser Gln Ala
 20 25 30
 Ile His Lys Tyr Asn Ala Tyr Arg Lys Ala Ala Ser Val Ile Ala Lys
 35 40 45
 Tyr Pro His Lys Ile Lys Ser Gly Ala Glu Ala Lys Lys Leu Pro Gly
 50 55 60
 Val Gly Thr Lys Ile Ala Glu Lys Ile Asp Glu Phe Leu Ala Thr Gly
 65 70 75 80
 Lys Leu Arg Lys Leu Glu Lys Ile Arg Gln Asp Asp Thr Ser Ser Ser
 85 90 95
 Ile Asn Phe Leu Thr Arg
 100

<210> 197

<211> 138

<212> PRT

<213> Homo sapien

<400> 197

Glu Ala Asn Glu Val Thr Asp Ser Ala Tyr Met Gly Ser Glu Ser Thr
 1 5 10 15
 Tyr Ser Glu Cys Glu Thr Phe Thr Asp Glu Asp Thr Ser Thr Leu Val
 20 25 30
 His Pro Glu Leu Gln Pro Glu Gly Asp Ala Asp Ser Ala Gly Gly Ser
 35 40 45
 Ala Val Pro Ser Glu Cys Leu Asp Ala Met Glu Glu Pro Asp His Gly
 50 55 60
 Ala Leu Leu Leu Leu Pro Gly Arg Pro His Pro His Gly Gln Ser Val
 65 70 75 80
 Ile Thr Val Ile Gly Gly Glu Glu His Phe Glu Asp Tyr Gly Glu Gly
 85 90 95
 Ser Glu Ala Glu Leu Ser Pro Glu Thr Leu Cys Asn Gly Gln Leu Gly
 100 105 110
 Cys Ser Asp Pro Ala Phe Leu Thr Pro Ser Pro Thr Lys Arg Leu Ser
 115 120 125

Ser Lys Lys Val Ala Arg Tyr Leu His Gln
130 135

<210> 198
<211> 100
<212> PRT
<213> Homo sapien

<400> 198
Met Gly Asp Val Lys Asn Phe Leu Tyr Ala Trp Cys Gly Lys Arg Lys
1 5 10 15
Met Thr Pro Ser Tyr Glu Ile Arg Ala Val Gly Asn Lys Asn Arg Gln
20 25 30
Lys Phe Met Cys Glu Val Gln Val Glu Gly Tyr Asn Tyr Thr Gly Met
35 40 45
Gly Asn Ser Thr Asn Lys Lys Asp Ala Gln Ser Asn Ala Ala Arg Asp
50 55 60
Phe Val Asn Tyr Leu Val Arg Ile Asn Glu Ile Lys Ser Glu Glu Val
65 70 75 80
Pro Ala Phe Gly Val Ala Ser Pro Pro Pro Leu Thr Asp Thr Pro Asp
85 90 95
Thr Thr Ala Asn
100

<210> 199
<211> 127
<212> PRT
<213> Homo sapien

<400> 199
Met Val Lys Glu Thr Thr Tyr Tyr Asp Val Leu Gly Val Lys Pro Asn
1 5 10 15
Ala Thr Gln Glu Leu Lys Lys Ala Tyr Arg Lys Leu Ala Leu Lys
20 25 30
Tyr His Pro Asp Lys Asn Pro Asn Glu Gly Glu Lys Phe Lys Gln Ile
35 40 45
Ser Gln Ala Tyr Glu Val Leu Ser Asp Ala Lys Lys Arg Glu Leu Tyr
50 55 60
Asp Lys Gly Gly Glu Gln Ala Ile Lys Glu Gly Gly Ala Gly Gly Gly
65 70 75 80
Phe Gly Ser Pro Met Asp Ile Phe Asp Met Phe Phe Gly Gly Gly Gly
85 90 95
Arg Met Gln Arg Glu Arg Arg Gly Lys Asn Val Val His Gln Leu Ser
100 105 110
Val Thr Leu Glu Asp Leu Tyr Asn Gly Ala Thr Arg Lys Leu Ala
115 120 125

<210> 200
<211> 90
<212> PRT
<213> Homo sapien

<400> 200
Met Ala Cys Pro Leu Asp Gln Ala Ile Gly Leu Leu Val Ala Ile Phe
1 5 10 15

His Lys Tyr Ser Gly Arg Glu Gly Asp Lys His Thr Leu Ser Lys Lys
 20 25 30
 Glu Leu Lys Glu Leu Ile Gln Lys Glu Leu Thr Ile Gly Ser Lys Leu
 35 40 45
 Gln Asp Ala Glu Ile Ala Arg Leu Met Glu Asp Leu Asp Arg Asn Lys
 50 55 60
 Asp Gln Glu Val Asn Phe Gln Glu Tyr Val Thr Phe Leu Gly Ala Leu
 65 70 75 80
 Ala Leu Ile Tyr Asn Glu Ala Leu Lys Gly
 85 90

<210> 201

<211> 120

<212> PRT

<213> Homo sapien

<400> 201

Met Glu Thr Pro Ser Gln Arg Arg Ala Thr Arg Ser Gly Ala Gln Ala
 1 5 10 15
 Ser Ser Thr Pro Leu Ser Pro Thr Arg Ile Thr Arg Leu Gln Glu Lys
 20 25 30
 Glu Asp Leu Gln Glu Leu Asn Asp Arg Leu Ala Val Tyr Ile Asp Arg
 35 40 45
 Val Arg Ser Leu Glu Thr Glu Asn Ala Gly Leu Arg Leu Arg Ile Thr
 50 55 60
 Glu Ser Glu Glu Val Val Ser Arg Glu Val Ser Gly Ile Lys Ala Ala
 65 70 75 80
 Tyr Glu Ala Glu Leu Gly Asp Ala Arg Lys Thr Leu Asp Ser Val Ala
 85 90 95
 Lys Glu Arg Ala Arg Leu Gln Leu Glu Leu Ser Lys Val Arg Glu Glu
 100 105 110
 Phe Lys Glu Leu Lys Ala Arg Asn
 115 120

<210> 202

<211> 177

<212> PRT

<213> Homo sapien

<400> 202

Met Ala Ala Gly Val Glu Ala Ala Ala Glu Val Ala Ala Thr Glu Ile
 1 5 10 15
 Lys Met Glu Glu Glu Ser Gly Ala Pro Gly Val Pro Ser Gly Asn Gly
 20 25 30
 Ala Pro Gly Pro Lys Gly Glu Gly Glu Arg Pro Ala Gln Asn Glu Lys
 35 40 45
 Arg Lys Glu Lys Asn Ile Lys Arg Gly Gly Asn Arg Phe Glu Pro Tyr
 50 55 60
 Ala Asn Pro Thr Lys Arg Tyr Arg Ala Phe Ile Thr Asn Ile Pro Phe
 65 70 75 80
 Asp Val Lys Trp Gln Ser Leu Lys Asp Leu Val Lys Glu Lys Val Gly
 85 90 95
 Glu Val Thr Tyr Val Glu Leu Leu Met Asp Ala Glu Gly Lys Ser Arg
 100 105 110
 Gly Cys Ala Val Val Glu Phe Lys Met Glu Glu Ser Met Lys Lys Ala

115 120 125
 Ala Glu Val Leu Asn Lys His Ser Leu Ser Gly Arg Pro Leu Lys Val
 130 135 140
 Lys Glu Asp Pro Asp Gly Glu His Ala Arg Arg Ala Met Gln Lys Ala
 145 150 155 160
 Gly Arg Leu Gly Ser Thr Val Phe Val Ala Asn Leu Asp Tyr Lys Val
 165 170 175
 Gly

<210> 203
 <211> 164
 <212> PRT
 <213> Homo sapien

<400> 203
 Met Arg Leu Ala Val Gly Ala Leu Leu Val Cys Ala Val Leu Gly Leu
 1 5 10 15
 Cys Leu Ala Val Pro Asp Lys Thr Val Arg Trp Cys Ala Val Ser Glu
 20 25 30
 His Glu Ala Thr Lys Cys Gln Ser Phe Arg Asp His Met Lys Ser Val
 35 40 45
 Ile Pro Ser Asp Gly Pro Ser Val Ala Cys Val Lys Lys Ala Ser Tyr
 50 55 60
 Leu Asp Cys Ile Arg Ala Ile Ala Ala Asn Glu Ala Asp Ala Val Thr
 65 70 75 80
 Leu Asp Ala Gly Leu Val Tyr Asp Ala Tyr Leu Ala Pro Asn Asn Leu
 85 90 95
 Lys Pro Val Val Ala Glu Phe Tyr Gly Ser Lys Glu Asp Pro Gln Thr
 100 105 110
 Phe Tyr Tyr Ala Val Ala Val Val Lys Lys Asp Ser Gly Phe Gln Met
 115 120 125
 Asn Gln Leu Arg Gly Lys Lys Ser Cys His Thr Gly Leu Gly Arg Ser
 130 135 140
 Ala Gly Trp Asn Ile Pro Ile Gly Leu Leu Tyr Cys Asp Leu Pro Glu
 145 150 155 160
 Pro Arg Lys Pro

<210> 204
 <211> 241
 <212> PRT
 <213> Homo sapien

<400> 204
 Met Ser Gly Glu Ser Ala Arg Ser Leu Gly Lys Gly Ser Ala Pro Pro
 1 5 10 15
 Gly Pro Val Pro Glu Gly Ser Ile Arg Ile Tyr Ser Met Arg Phe Cys
 20 25 30
 Pro Phe Ala Glu Arg Thr Arg Leu Val Leu Lys Ala Lys Gly Ile Arg
 35 40 45
 His Glu Val Ile Asn Ile Asn Leu Lys Asn Lys Pro Glu Trp Phe Phe
 50 55 60
 Lys Lys Asn Pro Phe Gly Leu Val Pro Val Leu Glu Asn Ser Gln Gly
 65 70 75 80

Gln Leu Ile Tyr Glu Ser Ala Ile Thr Cys Glu Tyr Leu Asp Glu Ala
 85 90 95
 Tyr Pro Gly Lys Lys Leu Leu Pro Asp Asp Pro Tyr Glu Lys Ala Cys
 100 105 110
 Gln Lys Met Ile Leu Glu Leu Phe Ser Lys Val Pro Ser Leu Val Gly
 115 120 125
 Ser Phe Ile Arg Ser Gln Asn Lys Glu Asp Tyr Asp Gly Leu Lys Glu
 130 135 140
 Glu Phe Arg Lys Glu Phe Thr Lys Leu Glu Glu Val Leu Thr Asn Lys
 145 150 155 160
 Lys Thr Thr Phe Phe Gly Gly Asn Ser Ile Ser Met Ile Asp Tyr Leu
 165 170 175
 Ile Trp Pro Trp Phe Glu Arg Leu Glu Ala Met Lys Leu Asn Glu Cys
 180 185 190
 Val Asp His Thr Pro Lys Leu Lys Leu Trp Met Ala Ala Met Lys Glu
 195 200 205
 Asp Pro Thr Val Ser Ala Leu Leu Thr Ser Glu Lys Asp Trp Gln Gly
 210 215 220
 Phe Leu Glu Leu Tyr Leu Gln Asn Ser Pro Glu Ala Cys Asp Tyr Gly
 225 230 235 240
 Leu

<210> 205

<211> 160

<212> PRT

<213> Homo sapien

<400> 205

Met Gln Ile Phe Val Lys Thr Leu Thr Gly Lys Thr Ile Thr Leu Glu
 1 5 10 15
 Val Glu Pro Ser Asp Thr Ile Glu Asn Val Lys Ala Lys Ile Gln Asp
 20 25 30
 Lys Glu Gly Ile Pro Pro Asp Gln Gln Arg Leu Ile Phe Ala Gly Lys
 35 40 45
 Gln Leu Glu Asp Gly Arg Thr Leu Ser Asp Tyr Asn Ile Gln Lys Glu
 50 55 60
 Ser Thr Leu His Leu Val Leu Arg Leu Arg Gly Gly Met Gln Ile Phe
 65 70 75 80
 Val Lys Thr Leu Thr Gly Lys Thr Ile Thr Leu Glu Val Glu Pro Ser
 85 90 95
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 115 120 125
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<212> PRT

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<400> 206

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 Phe Gly Pro Thr Gly Cys Gln Gly Ala Cys Leu Gly Cys Arg Asp His
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 Tyr Ser Gln Gln Ile Val Cys His Cys Arg Ala Gly Tyr Thr Gly Leu
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<212> PRT

<213> Homo sapien

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 35 40 45
 Ala Glu Ser Ala Trp Cys Tyr Leu Tyr Gly Thr Gly Ser Val Ala Gly
 50 55 60
 Val Tyr Leu Pro Gly Ser Arg Gln Thr Leu Ser Ile Tyr Gln Ala Leu
 65 70 75 80
 Lys Lys Gly Leu Leu Ser Ala Glu Val Ala Arg Leu Leu Leu Glu Ala
 85 90 95
 Gln Ala Ala Thr Gly Phe Leu Leu Asp Pro Val Lys Gly Glu Arg Leu
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<212> PRT

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 Arg Lys Glu Lys Asn Ile Lys Arg Gly Gly Asn Arg Phe Glu Pro Tyr
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 Glu Val Thr Tyr Val Glu Leu Leu Met Asp Ala Glu Gly Lys Ser Arg
 100 105 110
 Gly Cys Ala Val Val Glu Phe Lys Met Glu Glu Ser Met Lys Lys Ala
 115 120 125
 Ala Glu Val Leu Asn Lys His Ser Leu Ser Gly Arg Pro Leu Lys Val
 130 135 140
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 Ile

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<212> PRT

<213> Homo sapien

<400> 209

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 35 40 45
 Ala Glu Ser Asn His Ile Lys Leu Ser Gly Ser Asn Pro Tyr Thr Thr
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 65 70 75 80
 Val Pro Lys Arg Asp His Ala Leu Leu Glu Glu Gln Ser Lys Gln Gln
 85 90 95
 Ser Asn Glu His Leu Arg Arg Gln Phe Ala Ser Gln Ala Asn Val Val
 100 105 110
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 115 120 125

Glu Met Asn Gly Thr Leu Glu Asp Gln Leu Ser His Leu Lys Gln Tyr
 130 135 140
 Glu Arg Ser Ile Val Asp Tyr Lys Pro Asn Leu Asp Leu Leu Glu Gln
 145 150 155 160
 Gln His Gln Leu Ile Gln Glu Ala Leu Ile Phe Asp Asn Lys His Thr
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 Thr Ile Ala Arg
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<210> 210

<211> 156

<212> PRT

<213> Homo sapien

<400> 210

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 Trp Tyr Lys Gly Glu Arg Val Asp Gly Asn Ser Leu Ile Val Gly Tyr
 35 40 45
 Val Ile Gly Thr Gln Gln Ala Thr Pro Gly Pro Ala Tyr Ser Gly Arg
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 Glu Thr Ile Tyr Pro Asn Ala Ser Leu Leu Ile Gln Asn Val Thr Gln
 65 70 75 80
 Asn Asp Thr Gly Phe Tyr Thr Leu Gln Val Ile Lys Ser Asp Leu Val
 85 90 95
 Asn Glu Glu Ala Thr Gly Gln Phe His Val Tyr Pro Glu Leu Pro Lys
 100 105 110
 Pro Ser Ile Ser Ser Asn Asn Ser Asn Pro Val Glu Asp Lys Asp Ala
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<212> PRT

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 35 40 45
 Lys Glu Val Leu Leu Leu Val His Asn Leu Pro Gln His Leu Phe Gly
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85

90

<210> 212
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			20					25					30		
Leu	Gln	Glu	Glu	Val	Thr	Lys	Met	Asn	Leu	Leu	Asn	Gln	Gln	Ile	Gln
		35					40					45			
Glu	Glu	Leu	Ser	Arg	Val	Thr	Lys	Leu	Lys	Glu	Thr	Ala	Glu	Glu	Glu
	50					55					60				
Lys	Asp	Asp	Leu	Glu	Glu	Arg	Leu	Met	Asn	Gln	Leu	Ala	Glu	Leu	Asn
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				85				90					95		
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<210> 213
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<400> 213

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		20						25					30		
Leu	Ala	Ser	Tyr	Leu	Asp	Lys	Val	Arg	Ala	Leu	Glu	Ala	Ala	Asn	Gly
		35					40					45			
Glu	Leu	Glu	Val	Lys	Ile	Arg	Asp	Trp	Tyr	Gln	Lys	Gln	Gly	Pro	Gly
	50					55					60				
Pro	Ser	Arg	Asp	Tyr	Ser	His	Tyr	Tyr	Thr	Thr	Ile	Gln	Asp	Leu	Arg
65					70					75					80
Asp	Lys	Ile	Leu	Gly	Ala	Thr	Ile	Glu	Asn	Ser	Arg	Ile	Val	Leu	Gln
				85					90				95		
Ile	Asp	Asn	Ala	Arg	Leu	Ala	Ala	Asp	Asp	Phe	Arg	Thr	Lys	Phe	Glu
		100					105						110		
Thr	Glu	Gln	Ala	Leu	Arg	Met	Ser	Val	Glu	Ala	Asp	Ile	Asn	Gly	Leu
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	130					135						140			

<210> 214
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<213> Homo sapien

<400> 214

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      20           25           30
Asp Asn Gly Ala Lys Ser Val Val Leu Met Ser His Leu Gly Arg Pro
      35           40           45
Asp Gly Val Pro Met Pro Asp Lys Tyr Ser Leu Glu Pro Val Ala Val
      50           55           60
Glu Leu Arg Ser Leu Leu Gly Lys Asp Val Leu Phe Leu Lys Asp Cys
65           70           75           80
Val Gly Pro Glu Val Glu Lys Ala Cys Ala Asn Pro Ala Ala Gly Ser
      85           90           95
Val Ile Leu Leu Glu Asn Leu Arg Phe His Val Glu Glu Glu Gly Lys
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Gly Lys Asp Ala Ser Gly Asn Lys Val Lys Ala Glu Pro Ala Lys Ile
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<210> 215

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Leu Ala Leu Val Asp Val Leu Glu Asp Lys Leu Lys Gly Glu Met Met
      50           55           60
Asp Leu Gln His Gly Ser Leu Phe Leu Gln Thr Pro Lys Ile Val Ala
65           70           75           80
Asp Lys Asp Tyr Ser Val Thr Ala Asn Ser Lys Ile Val Val Val Thr
      85           90           95
Ala Gly Val Arg Gln Gln Glu Gly Glu Ser Arg Leu Asn Leu Val Gln
      100           105           110
Arg Asn Val Asn Val Phe Lys Phe Ile Ile Pro Gln Ile Val Lys Tyr
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Ser Pro Asp Cys Ile Ile Ile Val Val Ser Asn Pro Val Asp Ile Leu
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Thr Tyr Val Thr
145

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<210> 216

<211> 527

<212> PRT

<213> Homo sapien

<400> 216

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 35 40 45
 Pro Glu Ala Gly Glu Lys Val Leu Val Asn Gly Gly Leu Thr Pro Pro
 50 55 60
 Lys Ser Glu Asp Lys Val Ser Glu Asn Gly Gly Leu Arg Phe Pro Arg
 65 70 75 80
 Asn Thr Glu Arg Pro Pro Glu Thr Gly Pro Trp Arg Ala Pro Gly Pro
 85 90 95
 Trp Glu Lys Thr Pro Glu Ser Trp Gly Pro Ala Pro Thr Ile Gly Glu
 100 105 110
 Pro Ala Pro Glu Thr Ser Leu Glu Arg Ala Pro Ala Pro Ser Ala Val
 115 120 125
 Val Ser Ser Arg Asn Gly Gly Glu Thr Ala Pro Gly Pro Leu Gly Pro
 130 135 140
 Ala Pro Lys Asn Gly Thr Leu Glu Pro Gly Thr Glu Arg Arg Ala Pro
 145 150 155 160
 Glu Thr Gly Gly Ala Pro Arg Ala Pro Gly Ala Gly Arg Leu Asp Leu
 165 170 175
 Gly Ser Gly Gly Arg Ala Pro Val Gly Thr Gly Thr Ala Pro Gly Gly
 180 185 190
 Gly Pro Gly Ser Gly Val Asp Ala Lys Ala Gly Trp Val Asp Asn Thr
 195 200 205
 Arg Pro Gln Pro Pro Pro Pro Leu Pro Pro Pro Glu Ala Gln
 210 215 220
 Pro Arg Arg Leu Glu Pro Ala Pro Pro Arg Ala Arg Pro Glu Val Ala
 225 230 235 240
 Pro Glu Gly Glu Pro Gly Ala Pro Asp Ser Arg Ala Gly Gly Asp Thr
 245 250 255
 Ala Leu Ser Gly Asp Gly Asp Pro Pro Lys Pro Glu Arg Lys Gly Pro
 260 265 270
 Glu Met Pro Arg Leu Phe Leu Asp Leu Gly Pro Pro Gln Gly Asn Ser
 275 280 285
 Glu Gln Ile Lys Ala Arg Leu Ser Arg Leu Ser Leu Ala Leu Pro Pro
 290 295 300
 Leu Thr Leu Thr Pro Phe Pro Gly Pro Gly Pro Arg Arg Pro Pro Trp
 305 310 315 320
 Glu Gly Ala Asp Ala Gly Ala Ala Gly Gly Glu Ala Gly Gly Ala Gly
 325 330 335
 Ala Pro Gly Pro Ala Glu Glu Asp Gly Glu Asp Glu Asp Glu Asp Glu
 340 345 350
 Glu Glu Asp Glu Glu Ala Ala Ala Pro Gly Ala Ala Ala Gly Pro Arg
 355 360 365
 Gly Pro Gly Arg Ala Arg Ala Ala Pro Val Pro Val Val Val Ser Ser
 370 375 380
 Ala Asp Ala Asp Ala Ala Arg Pro Leu Arg Gly Leu Leu Lys Ser Pro
 385 390 395 400
 Arg Gly Ala Asp Glu Pro Glu Asp Ser Glu Leu Glu Arg Lys Arg Lys
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 Thr Pro Thr Asn Glu Leu Ser Val Gln Ala Pro Pro Glu Gly Asp Thr

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Pro	Gly	Asp	Gly	Phe	Pro	Ser	Asn	Asp	Ser	Gly	Phe	Gly	Gly	Ser	Phe
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Glu	Trp	Ala	Glu	Asp	Phe	Pro	Leu	Leu	Pro	Pro	Pro	Gly	Pro	Pro	Leu
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Cys	Phe	Ser	Arg	Phe	Ser	Val	Ser	Pro	Ala	Leu	Glu	Thr	Pro	Gly	Pro
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		515					520					525			



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C12N 15/12, A61K 38/17, C07K 14/47, 16/18, A61K 35/14		A3	(11) International Publication Number: WO 99/38973																					
			(43) International Publication Date: 5 August 1999 (05.08.99)																					
(21) International Application Number: PCT/US99/01642		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).																						
(22) International Filing Date: 26 January 1999 (26.01.99)																								
(30) Priority Data: <table border="0"> <tr> <td>09/015,029</td> <td>28 January 1998 (28.01.98)</td> <td>US</td> </tr> <tr> <td>09/015,022</td> <td>28 January 1998 (28.01.98)</td> <td>US</td> </tr> <tr> <td>09/040,828</td> <td>18 March 1998 (18.03.98)</td> <td>US</td> </tr> <tr> <td>09/040,831</td> <td>18 March 1998 (18.03.98)</td> <td>US</td> </tr> <tr> <td>09/122,192</td> <td>23 July 1998 (23.07.98)</td> <td>US</td> </tr> <tr> <td>09/122,191</td> <td>23 July 1998 (23.07.98)</td> <td>US</td> </tr> <tr> <td>09/219,245</td> <td>22 December 1998 (22.12.98)</td> <td>US</td> </tr> </table>		09/015,029	28 January 1998 (28.01.98)	US	09/015,022	28 January 1998 (28.01.98)	US	09/040,828	18 March 1998 (18.03.98)	US	09/040,831	18 March 1998 (18.03.98)	US	09/122,192	23 July 1998 (23.07.98)	US	09/122,191	23 July 1998 (23.07.98)	US	09/219,245	22 December 1998 (22.12.98)	US		
09/015,029	28 January 1998 (28.01.98)	US																						
09/015,022	28 January 1998 (28.01.98)	US																						
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09/122,192	23 July 1998 (23.07.98)	US																						
09/122,191	23 July 1998 (23.07.98)	US																						
09/219,245	22 December 1998 (22.12.98)	US																						
(71) Applicant: CORIXA CORPORATION [US/US]; Suite 200, 1124 Columbia Street, Seattle, WA 98104 (US).		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>																						
(72) Inventors: REED, Steven, G.; 2843 - 122nd Place N.E., Bellevue, WA 98005 (US). LODES, Michael, J.; 9223 - 36th Avenue S.W., Seattle, WA 98126 (US). FRUDAKIS, Tony, N.; P.O. Box 99232, Seattle, WA 99232-0232 (US). MOHAMATH, Raodoh; 4205 South Morgan, Seattle, WA 98118 (US).		(88) Date of publication of the international search report: 9 December 1999 (09.12.99)																						
(74) Agents: MAKI, David, J. et al.; Seed and Berry LLP, 6300 Columbia Center, 701 Fifth Avenue, Seattle, WA 98104-7092 (US).																								
(54) Title: COMPOUNDS FOR THERAPY AND DIAGNOSIS OF LUNG CANCER AND METHODS FOR THEIR USE																								
(57) Abstract Compounds and methods for treating lung cancer are provided. The inventive compounds include polypeptides containing at least a portion of a lung tumor protein. Vaccines and pharmaceutical compositions for immunotherapy of lung cancer comprising such polypeptides, or polynucleotides encoding such polypeptides, are also provided, together with polynucleotides for preparing the inventive polypeptides.																								

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 99/01642

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C12N15/12 A61K38/17 C07K14/47 C07K16/18 A61K35/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C12Q A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 96 30389 A (MILLENNIUM PHARMACEUTICALS, INC.; SHYJAN A.) 3 October 1996 see page 112 - page 127 ---	1-60
A	WO 96 02552 A (CYTOCLONYL PHARMACEUTICS, INC.; TORCZYNSKI R. ET AL.) 1 February 1996 see the whole document ---	1-60
A	YOU L ET AL.: "Identification of early growth response gene-1 (Egr-1) as a phorbol myristate-induced gene in lung cancer cells by differential mRNA display" AM. J. RESPIR. CELL MOL. BIOL., vol. 17, no. 5, November 1997, pages 617-624, XP002106654 see page 618, left-hand column, paragraph 3 --- -/--	1,2,4-7

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "A" document member of the same patent family

Date of the actual completion of the international search

21 June 1999

Date of mailing of the international search report

22 10. 1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
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Fax: (+31-70) 340-3016

Authorized officer

CUPIDO, M

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 16, 17, 24-26, 32, 33, 48-53 and 56-58 are directed to a method of treatment of the human/animal body the search has been carried out and based on the alleged effects of the composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see FURTHER INFORMATION sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

see FURTHER INFORMATION sheet, subject 1.

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/01642

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9630389 A	03-10-1996	US 5633161 A	27-05-1997
		AU 708746 B	12-08-1999
		AU 5437896 A	16-10-1996
		CA 2216717 A	03-10-1996
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